

FORM PTO-190 (REV. 11-2000)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER <b>GJE-65</b>
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			U.S. APPLICATION NO. (If known, see 37 CFR 1.5) <b>09/830807</b>
INTERNATIONAL APPLICATION NO. PCT/GB99/03721	INTERNATIONAL FILING DATE 09 Nov 1999	PRIORITY DATE CLAIMED 09 Nov 1998 (See #20 below)	
TITLE OF INVENTION Virulence Genes And Proteins, And Their Use			
APPLICANT(S) FOR DO/EO/US Helen Rachel Crooke, Enda Elizabeth Clarke, Paul Howard Everest, Gordon Dougan, David William Jacqueline Elizabeth Shea and Robert Graham Feldman			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:			
<p>1. <input checked="" type="checkbox"/> This is a <b>FIRST</b> submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a <b>SECOND</b> or <b>SUBSEQUENT</b> submission of items concerning a filing under 35 U.S.C. 371.</p> <p>3. <input type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.</p> <p>4. <input type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31).</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))</p> <p>a. <input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau).</p> <p>b. <input checked="" type="checkbox"/> has been communicated by the International Bureau.</p> <p>c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</p> <p>6. <input type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).</p> <p>a. <input type="checkbox"/> is attached hereto.</p> <p>b. <input type="checkbox"/> has been previously submitted under 35 U.S.C. 154(d)(4).</p> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))</p> <p>a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau).</p> <p>b. <input checked="" type="checkbox"/> have been communicated by the International Bureau.</p> <p>c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</p> <p>d. <input type="checkbox"/> have not been made and will not be made.</p> <p>8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)), <u>unsigned.</u></p> <p>10. <input type="checkbox"/> An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p>			
Items 11 to 20 below concern document(s) or information included:			
<p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input checked="" type="checkbox"/> A FIRST preliminary amendment.</p> <p>14. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>15. <input type="checkbox"/> A substitute specification.</p> <p>16. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>17. <input checked="" type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 37 CFR 1.821 - 1.825.</p> <p>18. <input type="checkbox"/> A second copy of the published international application under 35 U.S.C. 154(d)(4).</p> <p>19. <input type="checkbox"/> A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).</p> <p>20. <input checked="" type="checkbox"/> Other items or information:</p>			
Further priority dates: 17 December 1998; 13 January 1999; and 28 January 1999.			

APPLICATION NO. (if known) <b>09778350807</b>		INTERNATIONAL APPLICATION NO. PCT/GB99/03721		ATTORNEY'S DOCKET NUMBER <b>GJE-65</b>	
<b>21.</b> <input checked="" type="checkbox"/> The following fees are submitted: <b>BASIC NATIONAL FEE (37 CFR 1.492 (a) (1)-(5)):</b> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO ..... <b>\$1000.00</b>  International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO ..... <b>\$860.00</b>  International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO ..... <b>\$710.00</b>  International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) ..... <b>\$690.00</b> International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) ..... <b>\$100.00</b>  <b>ENTER APPROPRIATE BASIC FEE AMOUNT = \$860.00</b>				<b>CALCULATIONS    PTO USE ONLY</b>	
Surcharge of <b>\$130.00</b> for furnishing the oath or declaration later than months from the earliest claimed priority date (37 CFR 1.492(e)). <input type="checkbox"/> 20 <input type="checkbox"/> 30					
<b>CLAIMS</b>	<b>NUMBER FILED</b>	<b>NUMBER EXTRA</b>	<b>RATE</b>	<b>\$</b>	
Total claims	<u>22</u> - 20 =	<u>2</u>	x <b>\$18.00</b>	<b>\$36.00</b>	
Independent claims	<u>8</u> - 3 =	<u>5</u>	x <b>\$80.00</b>	<b>\$400.00</b>	
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ <b>\$270.00</b>	<b>\$0.00</b>	
<b>TOTAL OF ABOVE CALCULATIONS =</b>				<b>\$1,296.00</b>	
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.				<b>+ \$0.00</b>	
<b>SUBTOTAL =</b>				<b>\$1,296.00</b>	
Processing fee of <b>\$130.00</b> for furnishing the English translation later than months from the earliest claimed priority date (37 CFR 1.492(f)). <input type="checkbox"/> 20 <input type="checkbox"/> 30				<b>\$0.00</b>	
<b>TOTAL NATIONAL FEE =</b>				<b>\$1,296.00</b>	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). <b>\$40.00</b> per property +				<b>\$0.00</b>	
<b>TOTAL FEES ENCLOSED =</b>				<b>\$1,296.00</b>	
				<b>Amount to be refunded:</b>	<b>\$</b>
				<b>charged:</b>	<b>\$</b>
<p>a. <input type="checkbox"/> A check in the amount of \$ _____ to cover the above fees is enclosed.</p> <p>b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>19-0065</u> in the amount of \$ <u>1,296.00</u> to cover the above fees. A duplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>19-0065</u>. A duplicate copy of this sheet is enclosed.</p> <p>d. <input type="checkbox"/> Fees are to be charged to a credit card. <b>WARNING:</b> Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.</p>					
<p><b>NOTE:</b> Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.</p> <p>CORRESPONDENCE ADDRESS:</p>					
<b>CUSTOMER NUMBER</b>  <b>23,557</b>			<div style="text-align: center;">April 30, 2001</div> <div style="text-align: center;">DATE</div>		
			<div style="text-align: center;">SIGNATURE Glenn P. Ladwig</div> <div style="text-align: center;">NAME 46,853</div> <div style="text-align: center;">REGISTRATION NUMBER</div>		

09/830807

JC08 Rec'd PCT/PTO 30 APR 2001

April 30, 2001

Patent Application

Docket No. GJE-65

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants : Helen Rachel Crooke, Enda Elizabeth Clarke, Paul Howard Everest,  
Gordon Dougan, David William Holden, Jacqueline Elizabeth Shea,  
Robert Graham Feldman

Docket No. : GJE-241

For : Hydroxamic And Carboxylic Acid Derivatives having MMP and TNF  
Inhibitory Activity

PRELIMINARY AMENDMENT

Please amend the above-identified patent application as follows:

In the Specification

After page 17: Please insert as new page 18 the attached Abstract of the Disclosure.

In the claims

The following amendments are made with respect to the claims in the international application PCT/GB99/03721 attached as Annexes to the International Preliminary Examination Report (IPER). Therefore, please replace existing page 17 of the international application with the amended claim sheet (replacement page 17) of the annex attached to the IPER, and make the following amendments to the pending claims so that they read as follows:

Claim 1 (amended):

An isolated peptide encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 2 (amended):

The isolated peptide, according to claim 1, comprising an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

Claim 3 (amended):

An isolated polynucleotide which comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 4 (amended):

A host transformed to express a peptide encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a

homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 5 (amended):

A vaccine comprising a peptide, or the means for its expression, wherein said peptide is encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl1* to *msl16*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 6 (amended):

A vaccine comprising a microorganism having a virulence gene mutation, wherein the gene is selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl1* to *msl16*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 7 (amended):

The vaccine, according to claim 6, wherein said virulence gene mutation comprises a virulence gene deletion in two genes, wherein one gene encodes *tatA* and the other encodes *tatE*.

Claim 8 (amended):

The vaccine, according to claim 6, wherein the gene lies within a pathogenicity island.

Claim 9 (amended):

A method for screening potential drugs, or for the detection of virulence, wherein said method utilizes a peptide encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tataA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 10 (amended):

A method for treatment or prevention of a condition associated with infection by a Gram-negative bacterium, said method comprising administering a vaccine to a person or animal in need thereof, wherein said vaccine comprises a peptide, or a host transformed to express said peptide, wherein said peptide is encoded by an operon comprising a gene selected from the group consisting of *tataA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 11 (amended):

The method, according to claim 10, wherein the bacterium is *E. coli*.

Please add the following new claims:

12. The polynucleotide, according to claim 3, wherein said gene encodes a peptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

13. The host, according to claim 4, wherein said peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

14. The vaccine, according to claim 5, wherein said peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

15. The vaccine, according to claim 6, wherein said peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

16. The vaccine, according to claim 15, wherein said virulence gene mutation comprises a virulence gene deletion in two genes, wherein one gene encodes *tatA* and the other encodes *tatE*.

17. The vaccine, according to claim 15, wherein the gene lies within a pathogenicity island.

18. The method, according to claim 9, wherein said peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

19. The method, according to claim 9, wherein said peptide comprises an amino acid sequence as set forth in SEQ ID NO. 33.

20. The method, according to claim 10, wherein said peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

21. The method, according to claim 20, wherein the bacterium is *E. coli*.

22. A method for treatment or prevention of a condition associated with infection by a Gram-negative bacterium, said method comprising administering a nucleotide to a person or animal in need thereof, wherein said nucleotide comprises an operon including a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mid2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.




Remarks

Claims 1-11 have been amended and new claims 12-22 have been added.

No new matter has been added by these amendments.

The Commissioner is hereby authorized to charge any fees under 37 CFR 1.16 or 1.17 as required by this paper to Deposit Account 19-0065.

Respectfully Submitted



Glenn P. Ladwig

Patent Attorney

Registration No. 46,853

Phone No.: 352-375-8100

Address: Saliwanchik, Lloyd & Saliwanchik  
2421 N.W. 41<sup>st</sup> Street  
Suite A-1 Gainesville, FL 32606

GPL/la

Marked-up Version of Amended ClaimsClaim 1 (amended):

[A] An isolated peptide encoded by an operon, [including any of the genes identified herein as] wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to [16] *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, [having] wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof], for therapeutic use].

Claim 2 (amended):

[A] The isolated peptide, according to claim 1, comprising [any of the amino acid sequences identified herein as] an amino acid sequence selected from the group consisting of SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.

Claim 3 (amended):

[A] An isolated polynucleotide [encoding a peptide according to claim 1 or claim 2, for therapeutic use] which comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 4 (amended):

A host transformed to express a peptide [according to claim 1 or claim 2] encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 5 (amended):

A vaccine comprising a peptide [according to claim 1 or claim 2], or the means for its expression, wherein said peptide is encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 6 (amended):

A vaccine comprising a microorganism having a virulence gene mutation, wherein the gene is selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof [according to claim 1 or claim 2].

Claim 7 (amended):

[A] The vaccine, according to claim 6, [having] wherein said virulence gene mutation comprises a virulence gene deletion in two genes, wherein one gene encodes *tatA* and the other encodes *tatE*.

Claim 8 (amended):

[A] The vaccine, according to claim 6, wherein the gene lies within a pathogenicity island[, wherein the island comprises a gene identified herein].

Claim 9 (amended):

[Use of a product according to any of claims 1 to 4, or SEQ ID NO. 33,] A method for screening potential drugs, or for the detection of virulence, wherein said method utilizes a peptide encoded by an operon, wherein said operon comprises a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtl2*, and *ms1* to *ms16*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 10 (amended):

[Use of a product according to any of claims 1 to 4, for the manufacture of a medicament for use in the] A method for treatment or prevention of a condition associated with infection by a Gram-negative bacterium, said method comprising administering a vaccine to a person or animal in need thereof, wherein said vaccine comprises a peptide, or a host transformed to express said

peptide, wherein said peptide is encoded by an operon comprising a gene selected from the group consisting of *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2*, and *msl* to *msl6*, obtainable from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, wherein said homologue has at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof.

Claim 11 (amended):

[Use] The method, according to claim 10, wherein the bacterium is *E. coli*.

## VIRULENCE GENES AND PROTEINS, AND THEIR USE

### Field of the Invention

This invention relates to the identification of virulence genes and proteins, and their use. More particularly, it relates to their use in therapy and  
5 in screening for drugs.

### Background to the Invention

*E. coli* is a member of the *Enterobacteriaceae*, or enteric bacteria, which are Gram-negative microorganisms that populate the intestinal tracts of animals. Other members of this bacterial family include *Enterobacter*,  
10 *Klebsiella*, *Salmonella*, *Shigella* and *Yersinia*. Although *E. coli* is found normally in the human gastrointestinal tract, it has been implicated in human disease, including septicaemia, meningitis, urinary tract infection, wound infection, abscess formation, peritonitis and cholangitis.

The disease states caused by *E. coli* are dependent upon certain  
15 virulence determinants. For example, *E. coli* has been implicated in neonatal meningitis and a major determinant of virulence has been identified as the K1 antigen, which is a homopolymer of sialic acid. The K1 antigen may have a role in avoiding the host's immunological system and preventing phagocytosis.

### Summary of the Invention

20 The present invention is based on the identification of a series of virulence genes in *E. coli* K1, and also related organisms the products of which may be implicated in the pathogenicity of the organism.

According to one aspect of the present invention, a peptide is encoded by an operon including any of the genes identified herein as *mdoG*, *creC*, *recG*,  
25 *yggN*, *tatA*, *tatB*, *tatC*, *tatE*, *eck1*, *iroD*, *iroC*, *iroE*, *mtd2* and *ms1* to 16, from *E. coli* K1, or a homologue thereof in a Gram-negative bacterium, or a functional fragment thereof. Such a peptide is suitable for therapeutic use, e.g. when isolated.

The term "functional fragments" is used herein to define a part of the  
30 gene or peptide which retains similar therapeutic utility as the whole gene or peptide. For example, a functional fragment of the peptide may be used as an antigenic determinant, useful in a vaccine or in the production of antibodies.

A gene fragment may be used to encode the active peptide. Alternatively, the gene fragment may have utility in gene therapy, targetting the wild-type gene *in vivo* to exert a therapeutic effect.

A peptide according to the present invention may comprise any of the amino acid sequences identified herein as SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 23, 24, 25, 26, 28, 31, 29, 32 and 35-48.

The identification of these peptides as virulence determinants allows them to be used in a number of ways in the treatment of infection. For example, a host may be transformed to express a peptide according to the invention or modified to disrupt expression of the gene encoding the peptide. A vaccine may also comprise a peptide according to the invention, or the means for its expression, for the treatment of infection. In addition, a vaccine may comprise a microorganism having a virulence gene deletion, wherein the gene encodes a peptide according to the invention.

According to another aspect of the invention, the peptides or genes may be used for screening potential antimicrobial drugs or for the detection of virulence.

A further aspect of this invention is the use of any of the products identified herein, for the treatment or prevention of a condition associated with infection by a Gram-negative bacterium, in particular by *E. coli*.

#### Description of the Invention

The present invention has made use of signature-tagged mutagenesis (STM) (Hensel *et al*, Science, 1995;269:400-403) to screen *E. coli* K1 strain RS228 (Pluschke *et al*, Infection and Immunity 39:599-608) mini-Tn5 mutant bank for attenuated mutants, to identify virulence genes (and virulence determinants) of *E. coli*.

Although *E. coli* K1 was used as the microorganism to identify the virulence genes, corresponding genes in other enteric bacteria are considered to be within the scope of the present invention. For example, corresponding genes or encoded proteins may be found, based on sequence homology, in *Enterobacter*, *Klebsiella* and other genera implicated in human intestinal disease, including *Salmonella*, *Shigella* and *Yersinia*.

The term "virulence determinant" is used herein to define a product, e.g. a peptide or protein that may have a role in the maintenance of pathogenic bacteria. In particular, a virulence determinant is a bacterial protein or peptide that is implicated in the pathogenicity of the infectious or disease-causing microorganism.

A gene that encodes a virulence determinant may be termed a "virulence gene". Disruption of a virulence gene by way of mutation, deletion or insertion, will result in a reduced level of survival of the bacteria in a host, or a general reduction in the pathogenicity of the microorganism.

Signature-tagged mutagenesis has proved a very useful technique for identifying virulence genes, and their products. The technique relies on the ability of transposons to insert randomly into the genome of a microorganism, under permissive conditions. The transposons are individually marked for easy identification, and then introduced separately into a microorganism, resulting in disruption of the genome. Mutated microorganisms with reduced virulence are then detected by negative selection and the genes where insertional inactivation has occurred are identified and characterised.

A first stage in the STM process is the preparation of suitable transposons or transposon-like elements. A library of different transposons are prepared, each being incorporated into a vector or plasmid to facilitate transfer into the microorganism. The preparation of vectors with suitable transposons will be apparent to a skilled person in the art and is further disclosed in WO-A-96/17951. For the Gram-negative bacteria, e.g. *E. coli*, suitable transposons include Tn5 and Tn10. Having prepared the transposons, mutagenesis of a bacterial strain is then carried out to create a library of individually mutated bacteria.

Pools of the mutated microorganisms are then introduced into a suitable host. After a suitable length of time, the microorganisms are recovered from the host and those microorganisms that have survived in the host are identified, thereby also identifying the mutated strains that failed to survive, i.e. avirulent strains. Corresponding avirulent strains in a stored library are then used to identify the genes where insertional inactivation occurred. Usually, the site of



transposon insertion is identified by isolating the DNA flanking the transposons insertion site, and this permits characterisation of the genes implicated in virulence.

Once an avirulent microorganism has been identified, it is possible to determine more fully the potential role of the mutated gene in virulence, by infecting a suitable host animal with a lethal dose of the mutant. The survival time of the infected animal is compared with that of a control infected with the wild-type strain, and those animals surviving for longer periods than the control may be said to be infected with microorganisms having mutated virulence genes.

Alternatively, the potential role in virulence can be investigated by infecting an animal host with a mixture of the wild-type and mutant bacteria. After a suitable period of time, bacteria are harvested from organs of the host animal and the ratio of wild-type and mutant bacteria determined. This ratio is divided by the ratio of mutant to wild-type bacteria in the inoculum, to determine the competitive index (CI). Mutants which have a competitive index of less than 1 may be said to be avirulent.

It is possible that the gene which is inactivated by the insertion of the transposon may not be a true virulence gene, but may be having a polar effect on a downstream (virulence) gene. This can be determined by further experimentation, placing non-polar mutations in more defined regions of the gene, or mutating other adjacent genes, and establishing whether or not the mutant is avirulent.

Having characterised a virulence gene in *E. coli*, it is possible to use the gene sequence to establish homologies in other microorganisms. In this way it is possible to determine whether other microorganisms have similar virulence determinants. Sequence homologies may be established by searching in existing databases, e.g. EMBL or Genbank.

Virulence genes are often clustered together in distinct chromosomal regions called pathogenicity islands. Pathogenicity islands can be recognised as they are usually flanked by repeat sequences, insertion elements or tRNA genes. Also the G+C content is normally different from the remainder of the

chromosome, suggesting that they were acquired by horizontal transmission from another organism. For example the G+C content of the *E. coli* K12 genome is 52%. Any pathogenicity islands found in *E. coli* strains are likely to have a G+C content that varies from this average.

5       The identified virulence genes are likely to be useful both in generating attenuated vaccine strains and as a target for antimicrobials. The same may be true for homologues in Gram-negative bacteria in general.

10       For the purpose of this invention, the appropriate degree of homology is typically at least 30%, preferably at least 50%, 60% or 70%, and more preferably at least 80% or 90% (at the amino acid or nucleotide level).

15       Proteins according to the invention may be purified and isolated by methods known in the art. In particular, having identified the gene sequence, it will be possible to use recombinant techniques to express the genes in a suitable host. Active fragments and homologues can be identified and may be useful in therapy. For example, the proteins or their active fragments may be used as antigenic determinants in a vaccine, to elicit an immune response. They may also be used in the preparation of antibodies, for passive immunisation, or diagnostic applications. Suitable antibodies include monoclonal antibodies, or fragments thereof, including single chain fv  
20 fragments. Methods for the preparation of antibodies will be apparent to those skilled in the art.

25       The preparation of vaccines based on attenuated microorganisms is known to those skilled in the art. Vaccine compositions can be formulated with suitable carriers or adjuvants, e.g. alum, as necessary or desired, and used in therapy, to provide effective immunisation against *E. coli* or other Gram-negative bacteria. The preparation of vaccine formulations will be apparent to the skilled person.

30       More generally, and as is well known to those skilled in the art, a suitable amount of an active component of the invention can be selected, for therapeutic use, as can suitable carriers or excipients, and routes of administration. These factors will be chosen or determined according to known criteria such as the

nature/severity of the condition to be treated, the type or health of the subject etc.

The following Examples illustrate the invention. For the Examples, STM was used to screen an *E. coli* K1 mini-Tn5 mutant bank for attenuated mutants, using a mouse model of systemic infection. The basic procedure followed that disclosed in Hensel *et al, supra*. *E. coli* K1 containing a mini-Tn5 insertion within a virulence gene was not recovered from mice inoculated with a mixed population of mutants, and is therefore likely to be attenuated.

The DNA region flanking either side of the mini-Tn5 insertion was cloned by inverse PCR or by rescue of a kanamycin-resistance marker. In the latter case, chromosomal DNA from the STM-derived mutant was digested with restriction enzymes, ligated into the plasmid pUC19, and kanamycin-resistant clones selected after transformation into competent *E. coli* K12 cells. Subsequent cloning and sequencing was then performed and the gene sequences compared using sequences in publicly available sequence databases (EMBL) to help characterise the putative gene products.

#### Example 1

In a first mutant, two fragments of cloned DNA were sequenced. The nucleotide sequences are shown as SEQ ID NO. 1 and SEQ ID NO. 3 and a translated region of the DNA from SEQ ID NO. 1 is shown as SEQ ID NO. 2. SEQ ID NO. 1 shows 99.8% identity to the *mdoGH* region from *E. coli* K12 (EMBL database accession number AE000206) from nucleotides 2577 to 6908. This DNA fragment encodes the 5'-part of the *ymdD* gene, the entire *mdoG* gene and the 5'-part of the *mdoH* gene. The product of the *mdoG* gene is of unknown function, but is believed to be involved in the biosynthesis of membrane-derived oligosaccharides.

SEQ ID NO. 3 shows 98.3% identity to the 3'-part of the *mdoH* gene and downstream gene sequences from *E. coli* K12 (nucleotides 7187 to 7760). SEQ ID NO. 2 shows 99.6% identity to the *mdoG* protein from *E. coli* K12 (Swiss Prot accession number P33136) at amino acid 1 to 511.

The novel gene was tested for attenuation of virulence, using mixed infections, in a murine model of systemic infection (Achtman *et al.*, Infection and

Immunity, 1983; Vol. 39:315-335), and shown to be attenuated with a competitive index (CI) of 0.38. This confirms that the attenuation of the original transposon mutant is likely to be due to the disruption of the *mdoG* gene.

Polar and a non-polar deletion mutants of *mdoG* were constructed. The *mdoG* gene and flanking regions were amplified by PCR with oligonucleotides 5'-TGCTCTAGAGCCATTACTCAGAATGGG-3' (SEQ ID NO. 49) and 5'-CGCGAGCTCGACGACTGAATGATCCC-3' (SEQ ID NO. 50). The product was cloned into pUC19. A PCR product containing 5'- and 3'-terminal fragments of *mdoG* and the entire pUC19 sequence was then amplified by inverse PCR with the oligonucleotides 5'-TCCCCGGGTACTGCAGCACTCAACC-3' (SEQ ID NO. 51) and 5'-GATCCCGGGACCACTGAAATGCGTGC-3' (SEQ ID NO. 52). A non-polar kanamycin resistance cassette (*aphT*) was inserted in both orientations between the *mdoG* sequences to give a polar and a non-polar construct. The *mdoG::aphT* fusions were then transferred to the suicide vector pCDV442. The chromosomal copy of the *mdoG* was mutated by allelic transfer after conjugation of the pCDV442 constructs into wild type *E. coli* K1.

The constructed mutants were tested for attenuation of virulence in a murine model of systemic infection (Achtman et al., *supra*). Both the polar and the non-polar constructs were attenuated in virulence, with competitive indices of 0.37 and 0.35, respectively (mean CI from three mice each). This confirms that the attenuation of the original transposon mutant is likely to be due to the disruption of the *mdoG* gene.

#### Example 2

A second mutant was identified with a virulence gene having the nucleotide sequence shown in SEQ ID NO. 4 and the translated amino acid sequence shown as SEQ ID NO. 5. The mini-Tn5 transposon inserted at nucleotide 581 (SEQ ID NO. 4) and at amino acid 187 (SEQ ID NO. 5).

These sequences show 97.9% identity to the *creC* gene of *E. coli* K12 (EMBL and Genbank accession numbers M13608, AE000510 and U14003).

The *creC* protein from *E. coli* K12 belongs to the protein family of histidine kinases as well as to a protein family consisting of proteins containing a signal domain.

The novel gene was tested for attenuation of virulence (Achtman *et al*, *supra*), and shown to be attenuated with a competitive index of 0.09.

As the *E. coli* K12 *creC* gene is transcribed as part of an operon with the *creD* gene, it is possible that this attenuation is due to a polar effect on a presumed *E. coli* K1 *creD* gene.

#### Example 3

A third mutant had a nucleotide sequence shown as SEQ ID NO. 6 immediately following the mini-Tn5. A translation of this sequence is shown as SEQ ID NO. 7.

The nucleotide sequence shows 93.7% identity to the *recG* gene of *E. coli* K12, at nucleotides 5-146 (EMBL and Genbank accession numbers P24230 and M64367). This demonstrates that the disrupted gene is at least partially identical to the *recG* gene of *E. coli* K12. The *recG* gene of *E. coli* K12 encodes a 76.4kD protein which functions as ATP-dependent DNA helicase, and plays a critical role in DNA repair.

In tests for attenuation, the competitive index was shown to be 0.48. The *recG* gene is transcribed as the terminal gene of an operon, and it is therefore unlikely that this attenuation is due to a polar effect on another *E. coli* K1 gene.

#### Example 4

A fourth mutant had a transposon inserted within the nucleotide sequence shown as SEQ ID NO. 8, with a translation product shown as SEQ ID NO. 9.

The mini-Tn5 transposon inserted at nucleotide 359 and amino acid 80.

These sequences show 98.5% sequence identity to the *yggN* gene of *E. coli* K12 (EMBL accession number AE000378) at nucleotides 339-1054, and 99.6% identity at the amino acid level.

Although the sequence of the *yggN* gene is known, the function of its encoded protein has not yet been determined.

The novel gene was tested for attenuation of virulence, and shown to be attenuated with a competitive index of 0.43.

#### Example 5

Several mutants were also found with a transposon insertion within the same region. Cloning and sequencing the region revealed a nucleotide sequence shown as SEQ ID NO. 10. This sequence has homology with the *tatABCD* operon of *E. coli* K12 (EMBL and Genbank accession numbers AJ005830, AE000459 and AE000167). This operon encodes proteins of predicted mass 9.6 kD, 18.4 kD, 28.9 kD and 29.5 kD, which function as components of a Sec-independent protein export pathway. The pathway permits translocation of fully folded proteins to the periplasm through a gated pore, after the attachment of co-factors in the cytoplasm.

Translation of the nucleotide sequence revealed a protein corresponding to *tatA* (SEQ ID NO. 11), a sequence corresponding to *tatB* (SEQ ID NO. 12), a sequence corresponding to *tatC* (SEQ ID NO. 13) and a sequence corresponding to *tatD* (SEQ ID NO. 14).

The mini-Tn5 transposons in the mutants identified by STM are located at nucleotides 1429 and 2226 of SEQ ID NO. 10. These transposon insertions disrupt the *tatB* protein sequence at amino acid 50 and the *tatC* protein sequence at amino acid 143.

The *tatB* and *tatC* genes were tested for attenuation of virulence and were shown to be attenuated with competitive indices of 0.0012 and 0.0039, respectively. These genes were also attenuated in virulence when tested in single infections in the same model of systemic infection.

#### Example 6

A further mutant was insertionally inactivated within a region corresponding to the *tatE* gene of *E. coli* K12, shown as SEQ ID NO. 15. A translation of the sequence as shown as SEQ ID NO. 16. The *tatE* gene shows 98% identity to that of the *E. coli* K12 gene (accession number AE000167) at nucleotides 6719-7306.

To establish whether the *tatA*, *tatD* and *tatE* genes are required for virulence, non-polar deletion mutations were constructed in each. The regions of DNA flanking either side of the *tatA*, *tatD* and *tatE* genes were amplified with the following primers:

*tatA*

5'-TCG TCT AGA GAT GAT GGT GAT GGA GCG-3' (SEQ ID NO. 53)

5 5'-GAA CTG CAG CCA AAT ACT GAT ACC ACC C-3' (SEQ ID NO. 54)

5'-GAA CTG CAG GCT AAA ACA GAA GAC GCG-3' (SEQ ID NO. 55)

10 5'-CAT GCA TGC ACT CCA TAT GAC AAC CGC-3' (SEQ ID NO. 56)

Primers SEQ ID NO. 53 and SEQ ID NO. 54 were used to amplify DNA sequences upstream of *tatA*, Primers SEQ ID NO. 55 and SEQ ID NO. 56 were used to amplify DNA sequences downstream of *tatA*.

15 *tatD*

5'-TCG TCT AGA ATG AAG CTG CGC ATG AGG-3' (SEQ ID NO. 57)

20 5'-CAA CTG CAG TCG CAA ATT GCG AAC TGG-3' (SEQ ID NO. 58)

5'-CAA CTG CAG ACC GCA ACT TTT CGA CGC-3' (SEQ ID NO. 59)

5'-CAT GCA TGC CAG TGA GCC ATT GTT CCC-3' (SEQ ID NO. 60)

25 Primers SEQ ID NO. 57 and SEQ ID NO. 58 were used to amplify DNA sequences upstream of *tatD*, Primers SEQ ID NO. 59 and SEQ ID NO. 60 were used to amplify DNA sequences downstream of *tatD*.

*tatE*

30 5'-TGC TCT AGA TAC GAC TCT GAC AGG AGG-3' (SEQ ID NO. 61)

5'-TCA GAT ATC AAC TAC CAG CAG TTT GG-3' (SEQ ID NO. 62)

35 5'-TCA GAT ATC CAT AAA GAG TGA CGT GGC-3' (SEQ ID NO. 63)

5'-TGC TCT AGA AAA CGT GGC AAC AGA GCG-3' (SEQ ID NO. 64)

40 Primers SEQ ID NO. 61 and SEQ ID NO. 62 were used to amplify DNA sequences upstream of *tatE*, Primers SEQ ID NO. 63 and SEQ ID NO. 64 were used to amplify DNA sequences downstream of *tatE*.

After cloning these flanking DNA fragments into pUC19, a non-polar *aphT* kanamycin resistance cassette (Galan *et al.*, J.Bacteriol, 1992;174:4338-4349) was inserted between the flanking DNA fragments to replace the *tatA*, *tatD* and *tatE* genes. These DNA fragments were then transferred to the suicide  
5 vector pCVD442 (Blomfield *et. al.*, Mol. Micro., 1991;5:1447-1457). The chromosomal copies of the *E. coli* K1 *tatA*, *tatD* and *tatE* genes were then mutated by allelic transfer after conjugation of the pCVD442 constructs into wild type *E. coli* K1.

Disruptions of the *tatA*, *tatD* and *tatE* genes have been tested for  
10 attenuation of virulence (Achtman *et al.*, *supra*).

None of the genes was attenuated when deleted in isolation. The genes may still play a role in virulence, and to test this, mutants were prepared with deletions in both *tatA* and *tatE* genes. The double mutant was tested for attenuation in virulence using mixed infections with the wild-type strain and  
15 shown to be attenuated with a competitive index of 0.0017. It seems therefore that the *tatA*, *tatD* and *tatE* genes may be used in combination to create avirulent microorganisms.

Given the similarity of the *E. coli* K1 *tatABCD* genes to predicted *tatABCD* genes present in the *S. typhimurium* genome and *Neisseria meningitidis* genome it seemed likely that the *tat* system may also be required  
20 for virulence in these, and other, organisms. A deletion in the *S. typhimurium* *tatC* gene (SEQ ID NO. 17) was constructed by amplifying the DNA flanking either side of the *tatC* gene with the following primers:

25 5'-TGC TCT AGA AGG CGT TGT CGA TCC TG-3' (SEQ ID NO. 65)

5'-GAA CTG CAG GAA AAG GCC GAG CAG ACT G-3' (SEQ ID NO. 66)

5'-GAA CTG CAG TAC AGC CAT GTT TAC GGT-3' (SEQ ID NO. 67)

30 5'-CAT GCA TGC GGT GTA CGA CAG TTT GCG-3' (SEQ ID NO. 68)



Primers SEQ ID NO. 65 and SEQ ID NO. 66 were used to amplify DNA sequences downstream of the *S. typhimurium tatC* gene, Primers SEQ ID NO. 67 and SEQ ID NO. 68 were used to amplify DNA sequences upstream of the *S. typhimurium tatC* gene.

5       The encoded amino acid sequences for two regions of the *tatC* gene are shown as SEQ ID NO. 18 and SEQ ID NO. 19.

After cloning these flanking DNA fragments into pUC19, a non-polar kanamycin resistance cassette (*aphT*) was inserted between the flanking DNA fragments to replace the *S. typhimurium tatC* gene. This DNA fragment was then transferred to the suicide vector pCVD442. The chromosomal copy of the *S. typhimurium tatC* gene was then mutated by allelic transfer after conjugation of the pCVD442 construct into wild type *S. typhimurium* strains TML and SL1344.

10       The disrupted *S. typhimurium tatC* gene was tested for attenuation of virulence, using mixed and single infections in a murine model of systemic infection. For mixed infections, 6-7 week old *balbC* mice were inoculated intraperitoneally with  $10^4$  bacterial cells. Competitive indices were calculated after comparing the numbers of mutant and wild-type bacteria present in spleens after 3 days. For single infections, mice were inoculated either  
15       intraperitoneally or orally with varying doses and mouse survival monitored for 17 days. The strains were attenuated in virulence, the competitive indices of the SL1344 *tatC* and TML *tatC* deletion strains being 0.078 and 0.098, respectively.

20       In single infections, mouse survival was extended compared to the wild-type controls.

Sequence homology was also demonstrated with the *tat* sequence from *Neisseria meningitidis*. The gene sequence from *N. meningitidis* is shown as SEQ ID NO. 20 and the encoded amino acid sequence for *tatC* is shown as SEQ ID NO. 21.

30       To test for virulence, a deletion mutant was created using the following primers:

5'-TGCTCTAGACACATCATGGGCACACC-3' (SEQ ID NO. 69)

5'-GAACTGCAGAACCGTCCACATCAGGCG-3' (SEQ ID NO. 70)

5 5'-GAACTGCAGACCCTGCTTGCCATTCCG-3' (SEQ ID NO. 71)

5'-GAACTGCAGACCCTGCTTGCCATTCCG-3' (SEQ ID NO. 72)

10 Cloning of the DNA fragments and the *aphT* kanamycin resistance cassette into pUC19 followed the procedure outlined above for *S. typhimurium*. The chromosomal copy of the *N. meningitidis* *tatC* gene was mutated by transformation of the pUC19-based constructs into wild-type *N. meningitidis* cells.

15 Southern analysis of the resulting transformants indicated that all the transformants were merodiploids and contained both the wild-type and mutated copies of the *tatC* gene. This indicates that there is some selection against the isolation of mutants in which the *tatC* gene has been deleted.

Further studies on polar and non-polar constructs showed that transformants did not grow on selective media. This suggests that the *N. meningitidis* *tatC* gene is essential for the *in vitro* growth of this organism.

#### 20 Example 7

A further mutant was identified with a transposon insertion within a nucleotide sequence identified herein as SEQ ID NO. 22, at nucleotide 3981. The sequence defined herein as *eck1*, shows sequence homology to several 25 Group 1 glycosyltransferases from a number of bacteria. Sequence homology was also shown to the *gnd* gene of *E. coli* K12 (at nucleotides 4197-4604 of SEQ ID NO. 22).

The translation of the *E. coli* *eck1* gene is shown as SEQ ID NO. 26. The gene has been tested for attenuation of virulence, as described above, and 30 is shown to be attenuated with a competitive index of 0.025.

Several open reading frames (ORF) were also identified from the DNA sequence (SEQ ID NO. 22). The first of these is defined herein as MS1 and a translation product shown as SEQ ID NO. 25. The amino acid sequence is shown to have 50.3% identity to a putative glycosyl transferase from *E. coli*

serotype 0111 (TrEMBL database accession number AAD46732). The amino acid sequence also shows homology with the eck1 protein from *E. coli* K1 and also the TrsE protein from *Yersinia enterocolitica* (TrEMBL database accession number Q56917).

5 A second open reading frame identified herein as MS2 had the gene sequence shown as SEQ ID NO. 24. This shows sequence homology to the putative glycosyl transferase TrsC from *Yersinia enterocolitica* (TrRMBL database accession number Q56915), and also the glycosyl transferase WbnA from *E. coli* serotype 0113 (TrEMBL database accession number AAD50485).

10 A third open reading frame encodes a product identified herein as MS3 (SEQ ID NO. 23). The amino acid sequence shows 30.2% identity to a rhamnosyltransferase from *Streptococcus mutans*.

The gene sequence shown as SEQ ID NO. 22 may be at least part of a pathogenicity island, with multiple virulence genes being positioned in a cluster on the microorganism's genome.

#### 15 Example 8

A further mutant was identified having a transposon insertion within the *iroCDE* operon. The nucleotide sequences flanking either side of the mini-Tn5 insertion are shown as SEQ ID NO. 27 and SEQ ID NO. 30.

20 The mini-Tn5 transposon is inserted at nucleotide 1272 of SEQ ID NO. 27 and at nucleotide 1 of SEQ ID NO. 30, and interrupts the *iroD* gene. The N-terminal region of *iroD* is shown as SEQ ID NO. 29, and the C-terminal region is shown as SEQ ID NO. 31.

In addition to *iroD*, the gene shown as SEQ ID NO. 27 encodes a partial peptide with the amino acid sequence shown as SEQ ID NO. 28. This amino acid sequence shows 70.9% identity to the putative ATP binding cassette transporter *iroC* from *Salmonella typhi*.

25 The gene sequence shown as SEQ ID NO. 30 includes an open reading frame that encodes a peptide with the amino acid sequence shown as SEQ ID NO. 32 and this has sequence homology to the *iroE* protein from *Salmonella typhi*.

Testing the genes in a model for attenuation of virulence, as described above, showed that the *iroD* gene was attenuated with a competitive index of 0.107. The mini-Tn5 mutation in the *iroD* gene has been reintroduced into the wild-type *E. coli* K1 strain by P1 transduction. The resulting transductant is also  
5 attenuated in virulence with a competitive index of 0.1. This indicates that the attenuated phenotype is linked to the insertion within *iroD*. However, it is possible that the attenuation is due to a polar effect on the *E. coli* K1 *iroE* gene.

#### Example 9

A further mutant was identified with a transposon insertion within the  
10 nucleotide sequence shown as SEQ ID NO. 33. The transposon is inserted at nucleotide 2264 of SEQ ID NO. 33. The nucleotide sequence shows sequence homology to the *asI*A / *hemY* region of *E. coli* K12 (EMBL accession number AE000456). The *asI*A encodes an arylsulfatase homologue whereas *hemY* is involved in the biosynthesis of protoheme IX. This demonstrates that the  
15 disrupted region is at least partially identical to the *asI*A / *hemY* region of *E. coli* K12.

The transposon is inserted at nucleotide 2264 of SEQ ID NO. 33. This insertion site is 216 nucleotides downstream from the stop codon of the *hemY* gene and 472 nucleotides upstream from the start codon of the *asI*A gene.

The novel region has been tested for attenuation of virulence, as described above, and shown to be attenuated with a competitive index of 0.033. The mini-Tn5 mutation in this region has been reintroduced into the wild-type *E. coli* K1 strain by P1 transduction. The resulting transductant is also  
20 attenuated in virulence with a competitive index of 0.008. This indicates that the attenuated phenotype is linked to the transposon insertion in this region. However, polar and non-polar deletion mutants of *asI*A were constructed and tested for attenuation of virulence as described above.

Neither the polar nor the non-polar mutants were attenuated in virulence and this demonstrates that the attenuation of the original transposon mutant is  
30 not due to a polar effect on the *asI*A gene. This indicates that the transposon is disrupting some other function encoded within the intergenic region between *asI*A and *hemY*. For example there could be some untranslated RNA molecule,

such as a regulatory RNA similar to oxyS (Altuvia *et al.*, Cell, 1997;90:43-53), encoded within this region. Alternatively the transposon could be disrupting some DNA structure that may, for example, be involved in DNA replication. This DNA region is also present in the pathogen *Salmonella typhimurium* suggesting that it may be important for pathogenicity in other organisms. This region (SEQ ID NO. 33) may be used as a target, to identify anti-microbial drugs.

#### Example 10

A further mutant was identified and the DNA region flanking either side of the mini-Tn5 insertion was cloned and had the nucleotide sequence shown as SEQ ID NO. 34. This nucleotide sequence has homology with the *mtd2* gene of *Herpetosiphon aurantiacus* (EMBL accession number P25265), with the *mtd2* gene product functioning as a cytosine-specific methyltransferase. The *mtd2* gene is not found in the *E. coli* K12 genome and may represent a pathogenicity island.

The mini-Tn5 transposon insertions were located at nucleotides 4773 and 3764 of SEQ ID NO. 34 and were shown to interrupt the *mtd2* gene.

The amino acid sequence of the *mtd2* gene is shown as SEQ ID NO. 43.

The *E. coli* K1 *mtd2* gene was tested for attenuation of virulence, as described above, and shown to be attenuated with a competitive index of 0.073.

In addition to the *mtd2* gene, a series of open reading frames were also identified with translation products identified herein as MS4 to MS16, SEQ ID NOS. 48-44 and 42-35, respectively. As the open reading frames are located in a potential pathogenicity island, mutations in these genes may also result in attenuation in virulence. Further, since it is known that *E. coli* and other bacteria may encode peptides in different forms in the nucleotide sequence, the coding regions of some of these proteins may overlap. In addition, any amino acid sequence shown starting with Val may in fact start with Met.

CLAIMS

1. A peptide encoded by an operon including any of the genes identified herein as *tatA*, *tatB*, *tatC*, *tatE*, *mdoG*, *creC*, *recG*, *yggN*, *eck1*, *iroD*, *iroC*, *iroE*, *mta2* and *ms1* to 16, obtainable from *E. coli* K1, or a homologue thereof in a  
5 Gram-negative bacterium, having at least 30% homology at the amino acid or nucleotide level, or a functional fragment thereof, for therapeutic use.
2. A peptide according to claim 1, comprising any of the amino acid sequences identified herein as SEQ ID NOS. 2, 5, 7, 9, 11, 12, 13, 14, 16, 18, 19, 21, 23, 24, 25, 26, 28, 29, 31, 32 and 35-48.
- 10 3. A polynucleotide encoding a peptide according to claim 1 or claim 2, for therapeutic use.
4. A host transformed to express a peptide according to claim 1 or claim 2.
5. A vaccine comprising a peptide according to claim 1 or claim 2, or the means for its expression.
- 15 6. A vaccine comprising a microorganism having a virulence gene mutation, wherein the gene encodes a peptide according to claim 1 or claim 2.
7. A vaccine according to claim 6, having a virulence gene deletion in two genes, wherein one gene encodes *tatA* and the other encodes *tatE*.
8. A vaccine according to claim 6, wherein the gene lies within a  
20 pathogenicity island, wherein the island comprises a gene identified herein.
9. Use of a product according to any of claims 1 to 4, or SEQ ID NO. 33, for screening potential drugs or for the detection of virulence.
10. Use of a product according to any of claims 1 to 4, for the manufacture of a medicament for use in the treatment or prevention of a condition associated  
25 with infection by a Gram-negative bacterium.
11. Use according to claim 10, wherein the bacterium is *E. coli*.

The present invention is based on the identification of a series of virulence genes in *E. coli* K1, the products of which may be implicated in the pathogenicity of the organisms. The identification of the genes allows them, or their expressed products, to be used in a number of ways to treat infection.

DECLARATION AND POWER OF ATTORNEY

As a below-named inventor, I hereby declare that my residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of subject matter which is claimed and for which a patent is sought on an invention entitled  
**VIRULENCE GENES AND PROTEINS, AND THEIR USE**

the specification of which ☐ is attached hereto or

☒ was filed on 09 NOV 1999 as United States Application Number or PCT International Application Number PCT/GB99/03721 and was amended on 21 JAN 2001 (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56. I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for a patent or inventor's certificate, or PCT international application having a filing date before that of the application on which priority is claimed:

Prior Foreign Application Number(s)	Country	Foreign Filing Date	Priority Not Claimed	Certified Copy Attached? YES	NO
see attached sheet			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

As a named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:  
David R. Saliwanchik, Reg. 31,794; Jeff Lloyd, Reg. 35,589; Doran R. Pace, Reg. 38,261; Christine Q. McLeod, Reg. 36,213; Jay M. Sanders, Reg. 39,355; James S. Parker, Reg. 40,119 and Jean E. Kyle, Reg. 36,987; Frank C. Eisenschenk, Reg. 45,332; Seth M. Blum. Reg. 45,489

Direct all correspondence to:  
Saliwanchik, Lloyd & Saliwanchik  
2421 N.W. 41st Street, Suite A-1  
Gainesville, FL 32606-6669  
USA



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Country:	GB	Appln No:	9827815.3	Dated:	17 DEC 1998
Applicant:	Microscience Limited				
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Applicant:	Microscience Limited				

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C 1001 and that such willful false statements may jeopardise the validity of the application or any patent issued thereon.

1-00  
Full name of sole or First Inventor Helen Rachel CROOKE

Inventor's signature

Helen Croke

Residence address London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of Medicine at the Hammersmith Campus, Du Cane Road, London W12 ONN, United Kingdom

Country of Citizenship United Kingdom

Date of signature 20/4/01

2-00  
Full name of Second Inventor Enda Elizabeth CLARKE

Inventor's signature

Enda Clarke

Residence address London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of Medicine at the Hammersmith Campus, Du Cane Road, London W12 ONN, United Kingdom

Country of Citizenship United Kingdom

Date of signature 20-4-01

3-00  
Full name of Third Inventor Paul Howard EVEREST

Inventor's signature

Paul Everest

Residence address London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of Medicine at the Hammersmith Campus, Du Cane Road, London W12 ONN, United Kingdom

Country of Citizenship United Kingdom

Date of signature

30/4/2001

400 Full name of  
Fourth Inventor

Gordon DOUGAN

Inventor's signature



Residence address

London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of  
Medicine at the Hammersmith Campus, Du Cane Road, London  
W12 ONN, United Kingdom

Country of Citizenship

United Kingdom

Date of signature

8/5/01

500 Full name of  
Fifth Inventor

David William HOLDEN

Inventor's signature



Residence address

London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of  
Medicine at the Hammersmith Campus, Du Cane Road, London  
W12 ONN, United Kingdom

Country of Citizenship

United Kingdom

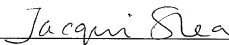
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600 Full name of  
Sixth Inventor

Jacqueline Elizabeth SHEA

Inventor's signature



Residence address

London, United Kingdom GBX

Post Office address

c/o Dept. of Infectious Diseases, Imperial College School of  
Medicine at the Hammersmith Campus, Du Cane Road, London  
W12 ONN, United Kingdom

Country of Citizenship

United Kingdom

Date of signature

29/5/01

7-00



Rafael

London, United Kingdom GBX

c/o Dept. of Infectious Diseases, Imperial College School of  
Medicine at the Hammersmith Campus, Du Cane Road, London  
W12 0NN, United Kingdom

Date of signature 25 APR 2001

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## SEQUENCE LISTING

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Glu Val Asp Gln Glu Tyr Val Thr Leu Lys Val Leu Asp Thr Gly Ser			
395	400	405	
ggg att cct gac tac gcg ctg tca cgt att ttt gaa cgc ttt tac tct			1299
Gly Ile Pro Asp Tyr Ala Leu Ser Arg Ile Phe Glu Arg Phe Tyr Ser			
410	415	420	425
ttg ccg cgt gca aat ggg caa aaa agc agc ggt ctg ggg ttg gcg ttt			1347
Leu Pro Arg Ala Asn Gly Gln Lys Ser Ser Gly Leu Gly Leu Ala Phe			
	430	435	440
gtc agt gag gtc gcc cgt ttg ttt aac ggc gaa gtc acg ctg cgc aac			1395
Val Ser Glu Val Ala Arg Leu Phe Asn Gly Glu Val Thr Leu Arg Asn			
	445	450	455
gtg cag gaa ggt ggc gtg ctg gcc tgc ctt cga ctt cac cgt cac ttc			1443
Val Gln Glu Gly Gly Val Leu Ala Ser Leu Arg Leu His Arg His Phe			
	460	465	470
aca tag cttcaaattc ttcccacata gtcttcgta			1478
Thr			
475			
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<211> 474			
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<213> Escherichia coli			
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Met Arg Ile Gly Met Arg Leu Leu Leu Gly Tyr Phe Leu Leu Val Ala			
1	5	10	15
Val Ala Ala Trp Phe Val Leu Ala Ile Phe Val Lys Glu Val Lys Pro			
	20	25	30
Gly Val Arg Arg Ala Thr Glu Gly Thr Leu Ile Asp Thr Ala Thr Leu			
	35	40	45
Leu Ala Glu Leu Ala Arg Pro Asp Leu Leu Ser Gly Asp Pro Thr His			
	50	55	60
Gly Gln Leu Ala Gln Ala Phe Asn Gln Leu Gln His Arg Pro Phe Arg			
	65	70	75
			80

Ala Asn Ile Gly Gly Ile Asn Lys Val Arg Asn Glu Tyr His Val Tyr  
85 90 95

Met Thr Asp Ala Gln Gly Lys Val Leu Phe Asp Ser Ala Asn Lys Ala  
100 105 110

Val Gly Gln Asp Tyr Ser Arg Trp Asn Asp Val Trp Leu Thr Leu Arg  
115 120 125

Gly Gln Tyr Gly Ala Arg Ser Thr Leu Gln Asn Pro Ala Asp Pro Glu  
130 135 140

Ser Ser Val Met Tyr Val Ala Ala Pro Ile Met Asp Gly Ser Arg Leu  
145 150 155 160

Ile Gly Val Leu Ser Val Gly Lys Pro Asn Ala Ala Met Ala Pro Val  
165 170 175

Ile Lys Arg Ser Glu Arg Arg Ile Leu Trp Ala Ser Ala Ile Leu Leu  
180 185 190

Gly Ile Ala Leu Val Ile Gly Ala Gly Met Val Trp Trp Ile Asn Arg  
195 200 205

Ser Ile Ala Arg Leu Thr Arg Tyr Ala Asp Ser Val Thr Asp Asn Lys  
210 215 220

Pro Val Pro Leu Pro Asp Leu Gly Ser Ser Glu Leu Arg Lys Leu Ala  
225 230 235 240

Gln Ala Leu Glu Ser Met Arg Val Lys Leu Glu Gly Lys Asn Tyr Ile  
245 250 255

Glu Gln Tyr Val Tyr Ala Leu Thr His Glu Leu Lys Ser Pro Leu Ala  
260 265 270

Ala Ile Arg Gly Ala Ala Glu Ile Leu Arg Glu Gly Pro Pro Pro Glu  
275 280 285

Val Val Ala Arg Phe Thr Asp Asn Ile Leu Thr Gln Asn Ala Arg Met  
290 295 300

Gln Ala Leu Val Glu Thr Leu Leu Arg Gln Ala Arg Leu Glu Asn Arg  
305 310 315 320

Gln Glu Val Val Leu Thr Ala Val Asp Val Ala Ala Leu Phe Arg Arg  
325 330 335



Val Ser Glu Ala Arg Thr Val Gln Leu Ala Glu Lys Asn Ile Thr Leu  
340 345 350

His Val Met Pro Thr Glu Val Asn Val Ala Ser Glu Pro Ala Leu Leu  
355 360 365

Glu Gln Ala Leu Gly Asn Leu Leu Asp Asn Ala Ile Asp Phe Thr Pro  
370 375 380

Glu Ser Gly Cys Ile Thr Leu Ser Ala Glu Val Asp Gln Glu Tyr Val  
385 390 395 400

Thr Leu Lys Val Leu Asp Thr Gly Ser Gly Ile Pro Asp Tyr Ala Leu  
405 410 415

Ser Arg Ile Phe Glu Arg Phe Tyr Ser Leu Pro Arg Ala Asn Gly Gln  
420 425 430

Lys Ser Ser Gly Leu Gly Leu Ala Phe Val Ser Glu Val Ala Arg Leu  
435 440 445

Phe Asn Gly Glu Val Thr Leu Arg Asn Val Gln Glu Gly Gly Val Leu  
450 455 460

Ala Ser Leu Arg Leu His Arg His Phe Thr  
465 470

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<211> 128

<212> DNA

<213> Escherichia coli

<220>

<221> CDS

<222> (1)..(126)

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atg aaa ggt cgc ctg tta gat gct gtc ccg ctc agt tcc cta acg ggc 48  
Met Lys Gly Arg Leu Leu Asp Ala Val Pro Leu Ser Ser Leu Thr Gly  
1 5 10 15

ggt ggc gca gcg ctt agt aac aag ctg gcg aaa atc aac ctg cat acc 96  
Val Gly Ala Ala Leu Ser Asn Lys Leu Ala Lys Ile Asn Leu His Thr  
20 25 30

gta cag gat tta ctc tta cac ctt cct ctg cg 128

Val Gln Asp Leu Leu Leu His Leu Pro Leu  
35 40

<210> 7

<211> 42

<212> PRT

<213> Escherichia coli

<400> 7

Met Lys Gly Arg Leu Leu Asp Ala Val Pro Leu Ser Ser Leu Thr Gly  
1 5 10 15

Val Gly Ala Ala Leu Ser Asn Lys Leu Ala Lys Ile Asn Leu His Thr  
20 25 30

Val Gln Asp Leu Leu Leu His Leu Pro Leu  
35 40

<210> 8

<211> 1174

<212> DNA

<213> Escherichia coli

<220>

<221> CDS

<222> (121)..(837)

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cacttttgta tcaatctggg gccagcaaat gctggcctga ttgttcttg aggggaagact 120

atg atg cgc aaa atg ctg ctg gcg gca gca ctt tca gtg acg gca atg 168  
Met Met Arg Lys Met Leu Leu Ala Ala Ala Leu Ser Val Thr Ala Met  
1 5 10 15

acc gct cac gcc gac tac cag tgc agc gtc acg ccg cgt gac gat gtg 216  
Thr Ala His Ala Asp Tyr Gln Cys Ser Val Thr Pro Arg Asp Asp Val  
20 25 30

att gtc agc ccg caa acc gtg cag gtg aag ggc gaa aac ggc aat ctg 264  
Ile Val Ser Pro Gln Thr Val Gln Val Lys Gly Glu Asn Gly Asn Leu  
35 40 45

gtg atc acg cca gac ggc aac gtg atg tat aac ggt aag caa tat tcc 312

Val	Ile	Thr	Pro	Asp	Gly	Asn	Val	Met	Tyr	Asn	Gly	Lys	Gln	Tyr	Ser		
	50					55					60						
ctg	aat	gcc	gcc	cag	cgc	gag	cag	gcg	aag	gat	tat	cag	gct	gaa	cta	360	
Leu	Asn	Ala	Ala	Gln	Arg	Glu	Gln	Ala	Lys	Asp	Tyr	Gln	Ala	Glu	Leu		
	65				70				75						80		
cgt	agc	acc	ctg	ccg	tgg	att	gat	gga	ggc	gcg	aaa	agc	cgc	gtc	gag	408	
Arg	Ser	Thr	Leu	Pro	Trp	Ile	Asp	Gly	Gly	Ala	Lys	Ser	Arg	Val	Glu		
				85				90						95			
aaa	gct	cgt	att	gcg	ctg	gat	aaa	att	atc	gtt	cag	gag	atg	ggc	gaa	456	
Lys	Ala	Arg	Ile	Ala	Leu	Asp	Lys	Ile	Ile	Val	Gln	Glu	Met	Gly	Glu		
		100					105						110				
agc	agc	aaa	atg	cgc	agc	cgt	ctg	acc	aaa	ctt	gat	gcg	cag	ctg	aaa	504	
Ser	Ser	Lys	Met	Arg	Ser	Arg	Leu	Thr	Lys	Leu	Asp	Ala	Gln	Leu	Lys		
		115					120					125					
gag	cag	atg	aac	cgc	att	atc	gaa	acg	cgc	agc	gat	ggc	ctg	acg	ttt	552	
Glu	Gln	Met	Asn	Arg	Ile	Ile	Glu	Thr	Arg	Ser	Asp	Gly	Leu	Thr	Phe		
	130					135					140						
cac	tat	aaa	gcc	att	gat	cag	gtt	cgt	gcc	gaa	ggc	cag	caa	tta	gtg	600	
His	Tyr	Lys	Ala	Ile	Asp	Gln	Val	Arg	Ala	Glu	Gly	Gln	Gln	Leu	Val		
	145					150				155				160			
aat	cag	gca	atg	ggc	gga	att	tta	cag	gac	agc	att	aat	gaa	atg	ggc	648	
Asn	Gln	Ala	Met	Gly	Gly	Ile	Leu	Gln	Asp	Ser	Ile	Asn	Glu	Met	Gly		
				165				170						175			
gcg	aaa	gcg	gtg	ctg	aaa	agc	ggc	ggt	aac	cca	tta	cag	aac	gtg	ctg	696	
Ala	Lys	Ala	Val	Leu	Lys	Ser	Gly	Gly	Asn	Pro	Leu	Gln	Asn	Val	Leu		
		180						185						190			
gga	agc	ctg	ggc	ggc	ctg	caa	tcc	tca	atc	caa	acc	gag	tgg	aaa	aag	744	
Gly	Ser	Leu	Gly	Gly	Leu	Gln	Ser	Ser	Ile	Gln	Thr	Glu	Trp	Lys	Lys		
		195					200						205				
cag	gaa	aaa	gat	ttc	cag	cag	ttt	ggc	aaa	gat	gtt	tgt	agc	cgc	gtt	792	
Gln	Glu	Lys	Asp	Phe	Gln	Gln	Phe	Gly	Lys	Asp	Val	Cys	Ser	Arg	Val		
		210					215				220						
gtg	act	ctg	gaa	gat	agc	cgc	aaa	gcc	ctg	gtc	ggg	aat	tta	aaa		837	
Val	Thr	Leu	Glu	Asp	Ser	Arg	Lys	Ala	Leu	Val	Gly	Asn	Leu	Lys			
	225					230				235							
taatcctcta ttttaagacg gcataatact tttttatgcc gtttaattct tcgttttgtt 897																	

acctgcctct aactttgtaa gggcgaattc tgcagatata catcacactg gcggccgctc 957  
 gagcatgcat cttaggggcc caattcgccc tatagtgagt cgtattacaa ttcactggcc 1017  
 gtcgttttac aaccgtctgt actgggaaaa ccttggcgtt acccaactta atcgcccttc 1077  
 agcacatccc ctttcgccca gctggcgtaa tagcgaaaag gcccgcacgg atcgcccttc 1137  
 caacagttgc gcacctgatg gccaatggac gcgcctg 1174

<210> 9

<211> 239

<212> PRT

<213> Escherichia coli

<400> 9

Met	Met	Arg	Lys	Met	Leu	Leu	Ala	Ala	Ala	Leu	Ser	Val	Thr	Ala	Met
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Thr	Ala	His	Ala	Asp	Tyr	Gln	Cys	Ser	Val	Thr	Pro	Arg	Asp	Asp	Val
			20					25						30	

Ile	Val	Ser	Pro	Gln	Thr	Val	Gln	Val	Lys	Gly	Glu	Asn	Gly	Asn	Leu
		35					40						45		

Val	Ile	Thr	Pro	Asp	Gly	Asn	Val	Met	Tyr	Asn	Gly	Lys	Gln	Tyr	Ser
	50					55					60				

Leu	Asn	Ala	Ala	Gln	Arg	Glu	Gln	Ala	Lys	Asp	Tyr	Gln	Ala	Glu	Leu
65						70				75				80	

Arg	Ser	Thr	Leu	Pro	Trp	Ile	Asp	Gly	Gly	Ala	Lys	Ser	Arg	Val	Glu
			85						90					95	

Lys	Ala	Arg	Ile	Ala	Leu	Asp	Lys	Ile	Ile	Val	Gln	Glu	Met	Gly	Glu
			100					105						110	

Ser	Ser	Lys	Met	Arg	Ser	Arg	Leu	Thr	Lys	Leu	Asp	Ala	Gln	Leu	Lys
		115					120					125			

Glu	Gln	Met	Asn	Arg	Ile	Ile	Glu	Thr	Arg	Ser	Asp	Gly	Leu	Thr	Phe
	130						135					140			

His	Tyr	Lys	Ala	Ile	Asp	Gln	Val	Arg	Ala	Glu	Gly	Gln	Gln	Leu	Val
145					150					155				160	

Asn Gln Ala Met Gly Gly Ile Leu Gln Asp Ser Ile Asn Glu Met Gly  
 165 170 175

Ala Lys Ala Val Leu Lys Ser Gly Gly Asn Pro Leu Gln Asn Val Leu  
 180 185 190

Gly Ser Leu Gly Gly Leu Gln Ser Ser Ile Gln Thr Glu Trp Lys Lys  
 195 200 205

Gln Glu Lys Asp Phe Gln Gln Phe Gly Lys Asp Val Cys Ser Arg Val  
 210 215 220

Val Thr Leu Glu Asp Ser Arg Lys Ala Leu Val Gly Asn Leu Lys  
 225 230 235

<210> 10

<211> 3406

<212> DNA

<213> Escherichia coli

<220>

<221> CDS

<222> (1007)..(1276)

<220>

<221> CDS

<222> (1280)..(1792)

<220>

<221> CDS

<222> (1798)..(2574)

<220>

<221> CDS

<222> (2604)..(3398)

<400> 10

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aaacggcacc aacatgaat tgctggcgga acgcggcggtg caggtgttct tcaactcaggt 120

ctttcgcgac agctttttcc atgctgatat gcaccctggc aacatcttcg taagctatga 180

acacccggaa aaccggaaat atatcgcat tgattgcggg attgttggt cgctaaacaa 240

agaagataaa cgctatctgg cggaaaactt tatcgccctt ttaaatcgcg actatcgcaa 300

18

His Asp Lys Glu Gln Val	Val Phe Asp Ile Gly Phe Ser Glu	
85	90	95
ctg cta ttg gtg ttc atc atc ggc ctc gtc gtt ctg ggg ccg caa cga		1351
Leu Leu Leu Val Phe Ile Ile Gly Leu Val Val Leu Gly Pro Gln Arg		
100	105	110
ctg cct gtg gcg gta aaa acg gta gcg ggc tgg att cgc gcg ttg cgt		1399
Leu Pro Val Ala Val Lys Thr Val Ala Gly Trp Ile Arg Ala Leu Arg		
115	120	130
tca ctg gcg aca acg gtg cag aac gaa ctg acc cag gag tta aaa ctc		1447
Ser Leu Ala Thr Thr Val Gln Asn Glu Leu Thr Gln Glu Leu Lys Leu		
	135	145
cag gag ttt cag gac agt ctg aaa aag gtt gaa aag gcg agc ctc act		1495
Gln Glu Phe Gln Asp Ser Leu Lys Lys Val Glu Lys Ala Ser Leu Thr		
	150	160
aac ctg acg ccc gaa ctg aaa gcg tcg atg gat gaa tta cgc cag gct		1543
Asn Leu Thr Pro Glu Leu Lys Ala Ser Met Asp Glu Leu Arg Gln Ala		
	165	175
gcg gag tcg atg aaa cgt tcc tac gtt gca aac gat cct gaa aag gcg		1591
Ala Glu Ser Met Lys Arg Ser Tyr Val Ala Asn Asp Pro Glu Lys Ala		
	180	190
agc gat gaa gcg cac acc atc cat aac ccg gtg gtg aaa gac aat gaa		1639
Ser Asp Glu Ala His Thr Ile His Asn Pro Val Val Lys Asp Asn Glu		
	195	210
act gcg cat gaa ggc gta acg cct gct gct gca caa acg cag gcc agt		1687
Thr Ala His Glu Gly Val Thr Pro Ala Ala Ala Gln Thr Gln Ala Ser		
	215	225
tcg ccg gaa cag aag cca gaa acc acg cca gag ccg gtg gta aaa cct		1735
Ser Pro Glu Gln Lys Pro Glu Thr Thr Pro Glu Pro Val Val Lys Pro		
	230	240
gct gcg gac gct gaa ccg aaa acc gct gca cct tcc cct tcg tcg agt		1783
Ala Ala Asp Ala Glu Pro Lys Thr Ala Ala Pro Ser Pro Ser Ser Ser		
	245	255
gat aaa ccg taaac atg tct gta gaa gat act caa ccg ctt atc acg cat		1833
Asp Lys Pro Met Ser Val Glu Asp Thr Gln Pro Leu Ile Thr His		
	260	270
ctg att gag ctg cgt aag cgt ctg ctg aac tgc att atc tcg gtg atc		1881

Leu Ile Glu Leu Arg Lys Arg Leu Leu Asn Cys Ile Ile Ser Val Ile	
275 280 285	
gtg ata ttc ctg tgt ctg gtc tat ttc gcc aat gac atc tat cac ctg	1929
Val Ile Phe Leu Cys Leu Val Tyr Phe Ala Asn Asp Ile Tyr His Leu	
290 295 300 305	
gta tcc gcg cca ctg atc aag cag ttg ccg caa ggt tca acg atg atc	1977
Val Ser Ala Pro Leu Ile Lys Gln Leu Pro Gln Gly Ser Thr Met Ile	
310 315 320	
gcc acc gac gtg gcc tcg ccg ttc ttt acg ccg atc aag ctg acc ttt	2025
Ala Thr Asp Val Ala Ser Pro Phe Phe Thr Pro Ile Lys Leu Thr Phe	
325 330 335	
atg gtg tcg ctg att ctg tca gcg ccg gtg att ctc tat cag gtg tgg	2073
Met Val Ser Leu Ile Leu Ser Ala Pro Val Ile Leu Tyr Gln Val Trp	
340 345 350	
gcg ttt atc gcc cca gcg ctg tat aag cat gaa cgt cgc ctg gtg gtg	2121
Ala Phe Ile Ala Pro Ala Leu Tyr Lys His Glu Arg Arg Leu Val Val	
355 360 365	
ccg ctg ctg gtt tcc agc tct ctg ctg ttt tat atc ggc atg gcg ttc	2169
Pro Leu Leu Val Ser Ser Ser Leu Leu Phe Tyr Ile Gly Met Ala Phe	
370 375 380 385	
gcc tac ttt gtg gtc ttt ccg ctg gca ttt ggc ttc ctt gcc aat acc	2217
Ala Tyr Phe Val Val Phe Pro Leu Ala Phe Gly Phe Leu Ala Asn Thr	
390 395 400	
gcg ccg gaa ggg gta cag gta tcc acc gac atc gcg agc tat tta agc	2265
Ala Pro Glu Gly Val Gln Val Ser Thr Asp Ile Ala Ser Tyr Leu Ser	
405 410 415	
ttc gtt atg gcg ctg ttt atg gcg ttt ggt gtc tcc ttt gaa gtg ccg	2313
Phe Val Met Ala Leu Phe Met Ala Phe Gly Val Ser Phe Glu Val Pro	
420 425 430	
gtg gca att gtg ctg ctg tgc tgg atg ggg att acc tcg cca gaa gac	2361
Val Ala Ile Val Leu Leu Cys Trp Met Gly Ile Thr Ser Pro Glu Asp	
435 440 445	
tta cgc aaa aaa cgc ccg tat gtg ctg gtt ggt gca ttc gtt gtc ggg	2409
Leu Arg Lys Lys Arg Pro Tyr Val Leu Val Gly Ala Phe Val Val Gly	
450 455 460 465	
atg ttg ctg acg ccg ccg gat gtc ttc tcg caa acg ctg ttg gcg atc	2457



Met Leu Leu Thr	Pro Pro Asp Val	Phe Ser Gln Thr	Leu Leu Ala Ile	
	470	475	480	
cct atg tac tgc ctg ttt gaa atc ggt gtc ttc ttc tca cgc ttt tac	2505			
Pro Met Tyr Cys Leu Phe Glu Ile Gly Val Phe Phe Ser Arg Phe Tyr				
	485	490	495	
gtt ggt aaa ggg cga aac cgg gaa gag gaa aac gac gct gaa gca gaa	2553			
Val Gly Lys Gly Arg Asn Arg Glu Glu Glu Asn Asp Ala Glu Ala Glu				
	500	505	510	
agc gaa aaa act gaa gaa taa attcaaccgc ccgtcagggc gggtgtcat atg	2606			
Ser Glu Lys Thr Glu Glu			Met	
	515	520		
gag tac agg atg ttt gat atc ggc gtt aat ttg acc agt tcg caa ttt	2654			
Glu Tyr Arg Met Phe Asp Ile Gly Val Asn Leu Thr Ser Ser Gln Phe				
	525	530	535	
gcg aaa gac cgt gat gat gtt gta gcg cgc gct ttt gac gcg gga gtt	2702			
Ala Lys Asp Arg Asp Asp Val Val Ala Arg Ala Phe Asp Ala Gly Val				
	540	545	550	
aat ggg cta ctc atc acc ggt acc aat ctg cgt gaa agc cag cag gcg	2750			
Asn Gly Leu Leu Ile Thr Gly Thr Asn Leu Arg Glu Ser Gln Gln Ala				
	555	560	565	
caa aag ctg gcg cgt cag tat tcg tcc tgt tgg tca acg gcg ggc gta	2798			
Gln Lys Leu Ala Arg Gln Tyr Ser Ser Cys Trp Ser Thr Ala Gly Val				
	570	575	580	585
cat cct cac gac agc agc cag tgg caa gct gtg act gaa gaa gcg att	2846			
His Pro His Asp Ser Ser Gln Trp Gln Ala Val Thr Glu Glu Ala Ile				
	590	595	600	
att gag ctg gcc gcg cag cca gaa gtg gtg gcg att ggt gaa tgt ggt	2894			
Ile Glu Leu Ala Ala Gln Pro Glu Val Val Ala Ile Gly Glu Cys Gly				
	605	610	615	
ctc gac ttt aac cgc aac ttt tcg acg ccg gaa gag cag gaa cgc gct	2942			
Leu Asp Phe Asn Arg Asn Phe Ser Thr Pro Glu Glu Gln Glu Arg Ala				
	620	625	630	
ttt gtt gcc cag cta cgc att gcc gca gaa tta aac atg ccg gta ttt	2990			
Phe Val Ala Gln Leu Arg Ile Ala Ala Glu Leu Asn Met Pro Val Phe				
	635	640	645	
atg cac tgt cgc gat gcc cac gag cgg ttt atg aca ttg ctg gag ccg	3038			

Met His Cys Arg Asp Ala His Glu Arg Phe Met Thr Leu Leu Glu Pro  
650 655 660 665

tgg ctg gat aaa ctg cct ggt gcg gtt ctt cat tgc ttt acc ggc aca 3086  
Trp Leu Asp Lys Leu Pro Gly Ala Val Leu His Cys Phe Thr Gly Thr  
670 675 680

cgc gaa gag atg cag gcg tgc gtg gcg tgt gga att tat atc ggc att 3134  
Arg Glu Glu Met Gln Ala Cys Val Ala Cys Gly Ile Tyr Ile Gly Ile  
685 690 695

acc ggt tgg gtt tgc gat gaa cga cgc ggg ctg gag ctg cgg gaa ttg 3182  
Thr Gly Trp Val Cys Asp Glu Arg Arg Gly Leu Glu Leu Arg Glu Leu  
700 705 710

ttg ccg ttg att ccg gcg gag aaa ttg ctg atc gaa act gat gcg ccg 3230  
Leu Pro Leu Ile Pro Ala Glu Lys Leu Leu Ile Glu Thr Asp Ala Pro  
715 720 725

tat ctg ctc cct cgc gat ctc acg cca aag cca tca tcc cgg cgc aac 3278  
Tyr Leu Leu Pro Arg Asp Leu Thr Pro Lys Pro Ser Ser Arg Arg Asn  
730 735 740 745

gag cca gcc cat ctg ccc cat att ttg caa cgt att gcg cac tgg cgt 3326  
Glu Pro Ala His Leu Pro His Ile Leu Gln Arg Ile Ala His Trp Arg  
750 755 760

gga gaa gat gcc gca tgg ctg gct gcc acc acg gat gcc aat gtc aaa 3374  
Gly Glu Asp Ala Ala Trp Leu Ala Ala Thr Thr Asp Ala Asn Val Lys  
765 770 775

aca ctg ttt ggg att gcg ttt tag agtttgcg 3406  
Thr Leu Phe Gly Ile Ala Phe  
780 785

&lt;210&gt; 11

&lt;211&gt; 89

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 11

Met Gly Gly Ile Ser Ile Trp Gln Leu Leu Ile Ile Ala Val Ile Val  
1 5 10 15

Val Leu Leu Phe Gly Thr Lys Lys Leu Gly Ser Ile Gly Ser Asp Leu  
20 25 30

Gly Ala Ser Ile Lys Gly Phe Lys Lys Ala Met Ser Asp Asp Glu Pro  
 35 40 45

Lys Gln Asp Lys Thr Ser Gln Asp Ala Asp Phe Thr Ala Lys Thr Ile  
 50 55 60

Ala Asp Lys Gln Ala Asp Thr Asn Gln Glu Gln Ala Lys Ile Glu Asp  
 65 70 75 80

Ala Lys Arg His Asp Lys Glu Gln Val  
 85

<210> 12

<211> 171

<212> PRT

<213> Escherichia coli

<400> 12

Val Phe Asp Ile Gly Phe Ser Glu Leu Leu Leu Val Phe Ile Ile Gly  
 1 5 10 15

Leu Val Val Leu Gly Pro Gln Arg Leu Pro Val Ala Val Lys Thr Val  
 20 25 30

Ala Gly Trp Ile Arg Ala Leu Arg Ser Leu Ala Thr Thr Val Gln Asn  
 35 40 45

Glu Leu Thr Gln Glu Leu Lys Leu Gln Glu Phe Gln Asp Ser Leu Lys  
 50 55 60

Lys Val Glu Lys Ala Ser Leu Thr Asn Leu Thr Pro Glu Leu Lys Ala  
 65 70 75 80

Ser Met Asp Glu Leu Arg Gln Ala Ala Glu Ser Met Lys Arg Ser Tyr  
 85 90 95

Val Ala Asn Asp Pro Glu Lys Ala Ser Asp Glu Ala His Thr Ile His  
 100 105 110

Asn Pro Val Val Lys Asp Asn Glu Thr Ala His Glu Gly Val Thr Pro  
 115 120 125

Ala Ala Ala Gln Thr Gln Ala Ser Ser Pro Glu Gln Lys Pro Glu Thr  
 130 135 140

Thr Pro Glu Pro Val Val Lys Pro Ala Ala Asp Ala Glu Pro Lys Thr  
 145 150 155 160

Ala Ala Pro Ser Pro Ser Ser Ser Asp Lys Pro  
165 170

<210> 13

<211> 258

<212> PRT

<213> Escherichia coli

<400> 13

Met Ser Val Glu Asp Thr Gln Pro Leu Ile Thr His Leu Ile Glu Leu  
1 5 10 15

Arg Lys Arg Leu Leu Asn Cys Ile Ile Ser Val Ile Val Ile Phe Leu  
20 25 30

Cys Leu Val Tyr Phe Ala Asn Asp Ile Tyr His Leu Val Ser Ala Pro  
35 40 45

Leu Ile Lys Gln Leu Pro Gln Gly Ser Thr Met Ile Ala Thr Asp Val  
50 55 60

Ala Ser Pro Phe Phe Thr Pro Ile Lys Leu Thr Phe Met Val Ser Leu  
65 70 75 80

Ile Leu Ser Ala Pro Val Ile Leu Tyr Gln Val Trp Ala Phe Ile Ala  
85 90 95

Pro Ala Leu Tyr Lys His Glu Arg Arg Leu Val Val Pro Leu Leu Val  
100 105 110

Ser Ser Ser Leu Leu Phe Tyr Ile Gly Met Ala Phe Ala Tyr Phe Val  
115 120 125

Val Phe Pro Leu Ala Phe Gly Phe Leu Ala Asn Thr Ala Pro Glu Gly  
130 135 140

Val Gln Val Ser Thr Asp Ile Ala Ser Tyr Leu Ser Phe Val Met Ala  
145 150 155 160

Leu Phe Met Ala Phe Gly Val Ser Phe Glu Val Pro Val Ala Ile Val  
165 170 175

Leu Leu Cys Trp Met Gly Ile Thr Ser Pro Glu Asp Leu Arg Lys Lys  
180 185 190

Arg Pro Tyr Val Leu Val Gly Ala Phe Val Val Gly Met Leu Leu Thr

195

200

205

Pro Pro Asp Val Phe Ser Gln Thr Leu Leu Ala Ile Pro Met Tyr Cys  
 210 215 220

Leu Phe Glu Ile Gly Val Phe Phe Ser Arg Phe Tyr Val Gly Lys Gly  
 225 230 235 240

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Glu Glu

<210> 14

<211> 264

<212> PRT

<213> Escherichia coli

<400> 14

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Val Asn Gly Leu Leu Ile Thr Gly Thr Asn Leu Arg Glu Ser Gln Gln  
 35 40 45

Ala Gln Lys Leu Ala Arg Gln Tyr Ser Ser Cys Trp Ser Thr Ala Gly  
 50 55 60

Val His Pro His Asp Ser Ser Gln Trp Gln Ala Val Thr Glu Glu Ala  
 65 70 75 80

Ile Ile Glu Leu Ala Ala Gln Pro Glu Val Val Ala Ile Gly Glu Cys  
 85 90 95

Gly Leu Asp Phe Asn Arg Asn Phe Ser Thr Pro Glu Glu Gln Glu Arg  
 100 105 110

Ala Phe Val Ala Gln Leu Arg Ile Ala Ala Glu Leu Asn Met Pro Val  
 115 120 125

Phe Met His Cys Arg Asp Ala His Glu Arg Phe Met Thr Leu Leu Glu  
 130 135 140

Pro Trp Leu Asp Lys Leu Pro Gly Ala Val Leu His Cys Phe Thr Gly

145 150 155 160

Thr Arg Glu Glu Met Gln Ala Cys Val Ala Cys Gly Ile Tyr Ile Gly  
165 170 175

Ile Thr Gly Trp Val Cys Asp Glu Arg Arg Gly Leu Glu Leu Arg Glu  
180 185 190

Leu Leu Pro Leu Ile Pro Ala Glu Lys Leu Leu Ile Glu Thr Asp Ala  
195 200 205

Pro Tyr Leu Leu Pro Arg Asp Leu Thr Pro Lys Pro Ser Ser Arg Arg  
210 215 220

Asn Glu Pro Ala His Leu Pro His Ile Leu Gln Arg Ile Ala His Trp  
225 230 235 240

Arg Gly Glu Asp Ala Ala Trp Leu Ala Ala Thr Thr Asp Ala Asn Val  
245 250 255

Lys Thr Leu Phe Gly Ile Ala Phe  
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<210> 15

<211> 586

<212> DNA

<213> Escherichia coli

<220>

<221> CDS

<222> (170)..(370)

<400> 15

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tgattcacct tgttacagat tgctattgtg tgcgcgcgctc gaatgaccgt taatattctc 120

tggttttttaa ggcgcgttct gttgccggtt atatgtcaag aaggtatct atg ggt gag 178  
Met Gly Glu

1

att agt att acc aaa ctg ctg gta gtt gcg gcg ctg gtc gtt ctg ctg 226  
Ile Ser Ile Thr Lys Leu Leu Val Val Ala Ala Leu Val Val Leu Leu

5

10

15

ttt ggg act aag aag tta cgt acg ctg ggc gga gac ctt gga gcg gcc 274

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Phe Gly Thr Lys Lys Leu Arg Thr Leu Gly Gly Asp Leu Gly Ala Ala
20                      25                      30                      35

att aaa ggg ttc aag aag gcg atg aat gat gac gat gct gcg gcg aaa 322
Ile Lys Gly Phe Lys Lys Ala Met Asn Asp Asp Ala Ala Ala Lys
40                      45                      50

aaa ggc gca gac gtt gat ctt cag gct gaa aag ctc tct cat aaa gag 370
Lys Gly Ala Asp Val Asp Leu Gln Ala Glu Lys Leu Ser His Lys Glu
55                      60                      65

tgacgtggcg agcaggacgc tccctcaata tcttgctcga tacaaaaacc cgcttcaaaa 430

agcggtgttt ttatcagaca gatgtaagta attattacag gattacttaa cttccatccc 490

tttcgcctgc aaatcggcgt ggtaagaaga gcggacaaac ggaccgcatg cagcatgggt 550

aaagcccatc gccagcgctt cgctttcatt tcgctg 586

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&lt;210&gt; 16

&lt;211&gt; 67

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 16

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Met Gly Glu Ile Ser Ile Thr Lys Leu Leu Val Val Ala Ala Leu Val
1                      5                      10                      15

Val Leu Leu Phe Gly Thr Lys Lys Leu Arg Thr Leu Gly Gly Asp Leu
20                      25                      30

Gly Ala Ala Ile Lys Gly Phe Lys Lys Ala Met Asn Asp Asp Asp Ala
35                      40                      45

Ala Ala Lys Lys Gly Ala Asp Val Asp Leu Gln Ala Glu Lys Leu Ser
50                      55                      60

His Lys Glu
65

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&lt;210&gt; 17

&lt;211&gt; 4200

&lt;212&gt; DNA

&lt;213&gt; Salmonella typhimurium

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (947)..(1444)

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1450)..(1722)

&lt;400&gt; 17

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atcacagagg aacatgtatg ggtggtatca gtatttgca gttgttgatt gttgccgtta 180
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cggagtcgat gaaacgcacc tacagcgcta acgatccga acaagcgagc gatgaagcgc 720
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cgctgttgt cgaattctcc cctcgtgta gtgataaacc gtaaac atg gct gta 955
                                     Met Ala Val
                                     1
gaa gat act caa ccg ctt atc acg cat ctg atc gag ttg cgt aag cgc 1003
Glu Asp Thr Gln Pro Leu Ile Thr His Leu Ile Glu Leu Arg Lys Arg
      5              10              15
ctg cta aac tgc atc gtc gca gta ctt ctg att ttt ctg gcg tta att 1051

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Leu Leu Asn Cys Ile Val Ala Val Leu Leu Ile Phe Leu Ala Leu Ile	
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Tyr Phe Ala Asn Asp Ile Tyr His Leu Val Ala Ala Pro Leu Ile Lys	
40 45 50	
cag atg ccg caa ggg gcg aca atg att gcg acg gat gtg gcg tcg ccg	1147
Gln Met Pro Gln Gly Ala Thr Met Ile Ala Thr Asp Val Ala Ser Pro	
55 60 65	
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Phe Phe Thr Pro Ile Lys Leu Thr Phe Met Val Ser Leu Ile Leu Ser	
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gcg cct gtc att ttg tac cag gtt tgg gcc ttt atc gcc ccg gcg ctg	1243
Ala Pro Val Ile Leu Tyr Gln Val Trp Ala Phe Ile Ala Pro Ala Leu	
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Tyr Lys His Glu Arg Arg Leu Val Val Pro Leu Leu Val Ser Ser Ser	
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ctg ctt ttc tat att ggt atg gcc ttc gcc tat ttt gtc gta ttc cct	1339
Leu Leu Phe Tyr Ile Gly Met Ala Phe Ala Tyr Phe Val Val Phe Pro	
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Leu Ala Phe Gly Phe Leu Thr His Thr Ala Pro Glu Gly Val Gln Val	
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Ser Thr Asp Ile Ala Ser Tyr Leu Ser Phe Val Met Ala Leu Phe Met	
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Ala Phe Ala Phe Glu Val Pro Val Ala Ile Val Leu Leu Cys Trp	
165 170 175	
atg ggc atc acc acg cca gaa gat ttg cgt aaa aaa ccg cct tat atc	1533
Met Gly Ile Thr Thr Pro Glu Asp Leu Arg Lys Lys Arg Pro Tyr Ile	
180 185 190	
ctg gtc ggg gca ttc att gtg gga atg ctg ctt acg ccg cca gat gtt	1581
Leu Val Gly Ala Phe Ile Val Gly Met Leu Leu Thr Pro Pro Asp Val	
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Phe Ser Gln Thr Leu Leu Ala Ile Pro Met Tyr Cys Leu Phe Glu Ile  
 215 220 225  
 ggc gtt ttc tgc tca cgc ttt tat gtc ggt aag cga cgg acg cgc gac 1677  
 Gly Val Phe Cys Ser Arg Phe Tyr Val Gly Lys Arg Arg Thr Arg Asp  
 230 235 240  
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&lt;210&gt; 18

&lt;211&gt; 166

&lt;212&gt; PRT

<213> *Salmonella typhimurium*

&lt;400&gt; 18

Met Ala Val Glu Asp Thr Gln Pro Leu Ile Thr His Leu Ile Glu Leu  
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Arg Lys Arg Leu Leu Asn Cys Ile Val Ala Val Leu Leu Ile Phe Leu  
 20 25 30

Ala Leu Ile Tyr Phe Ala Asn Asp Ile Tyr His Leu Val Ala Ala Pro  
 35 40 45

Leu Ile Lys Gln Met Pro Gln Gly Ala Thr Met Ile Ala Thr Asp Val  
 50 55 60

Ala Ser Pro Phe Phe Thr Pro Ile Lys Leu Thr Phe Met Val Ser Leu  
 65 70 75 80

Ile Leu Ser Ala Pro Val Ile Leu Tyr Gln Val Trp Ala Phe Ile Ala  
 85 90 95

Pro Ala Leu Tyr Lys His Glu Arg Arg Leu Val Val Pro Leu Leu Val  
 100 105 110

Ser Ser Ser Leu Leu Phe Tyr Ile Gly Met Ala Phe Ala Tyr Phe Val  
 115 120 125

Val Phe Pro Leu Ala Phe Gly Phe Leu Thr His Thr Ala Pro Glu Gly  
 130 135 140

Val Gln Val Ser Thr Asp Ile Ala Ser Tyr Leu Ser Phe Val Met Ala  
 145 150 155 160

Leu Phe Met Ala Phe Ala  
 165

&lt;210&gt; 19

&lt;211&gt; 91

&lt;212&gt; PRT

<213> *Salmonella typhimurium*

&lt;400&gt; 19

Phe Glu Val Pro Val Ala Ile Val Leu Leu Cys Trp Met Gly Ile Thr  
 1 5 10 15

Thr	Pro	Glu	Asp	Leu	Arg	Lys	Lys	Arg	Pro	Tyr	Ile	Leu	Val	Gly	Ala
				20				25					30		
Phe	Ile	Val	Gly	Met	Leu	Leu	Thr	Pro	Pro	Asp	Val	Phe	Ser	Gln	Thr
				35			40					45			
Leu	Leu	Ala	Ile	Pro	Met	Tyr	Cys	Leu	Phe	Glu	Ile	Gly	Val	Phe	Cys
		50				55				60					
Ser	Arg	Phe	Tyr	Val	Gly	Lys	Arg	Arg	Thr	Arg	Asp	Glu	Asp	Asn	Glu
	65				70				75					80	
Ala	Glu	Thr	Glu	Lys	Ala	Glu	His	Thr	Glu	Asp					
				85				90							

&lt;210&gt; 20

&lt;211&gt; 2601

&lt;212&gt; DNA

&lt;213&gt; Neisseria meningitidis

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1572)..(2339)

&lt;400&gt; 20

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 ttgtgcgggc ttggccgaaa aagccgctcg cacagaatct tgaattata ttaaaatccg 240  
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 aactgtatgt tctgcaaaat cgccgcaaaa gacattccgg cgaaaccgt ctatgaagac 360  
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 Val Ser Glu Thr Gln Asn Glu Gln Pro Val Gln Pro Leu  
 1 5 10  
 gtc gag cat ctc atc gag ctg cgc cgc cgc ctg atg tgg acg gtt gtc 1658  
 Val Glu His Leu Ile Glu Leu Arg Arg Arg Leu Met Trp Thr Val Val  
 15 20 25  
 ggt atc tta gtc tgc ttt ttc ggc cta atg ccg ttt gcc caa caa ctc 1706  
 Gly Ile Leu Val Cys Phe Phe Gly Leu Met Pro Phe Ala Gln Gln Leu  
 30 35 40 45  
 tat act ttt atc gcc gac ccg ctg atg gca aac ctg ccc aaa gac acc 1754  
 Tyr Thr Phe Ile Ala Asp Pro Leu Met Ala Asn Leu Pro Lys Asp Thr  
 50 55 60

agc atg att gcc acc gat gtc atc gca cca ttt ttc gtg ccg gtc aaa 1802  
 Ser Met Ile Ala Thr Asp Val Ile Ala Pro Phe Phe Val Pro Val Lys  
 65 70 75

gtt acc ctg atg gcg gca ttt tta att tcg ctg ccg cat acg ctc tac 1850  
 Val Thr Leu Met Ala Ala Phe Leu Ile Ser Leu Pro His Thr Leu Tyr  
 80 85 90

caa atc tgg gca ttc gtc gcc ccc gca ctc tac caa aac gaa aaa cgc 1898  
 Gln Ile Trp Ala Phe Val Ala Pro Ala Leu Tyr Gln Asn Glu Lys Arg  
 95 100 105

ctg att acg ccg ctc gtc ctc tcc agc gtc agc ctg ttt ttc atc ggc 1946  
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atg gca ttt gcc tac ttt ttg gtt ttc ccc gtc att ttc aaa ttc ctt 1994  
 Met Ala Phe Ala Tyr Phe Leu Val Phe Pro Val Ile Phe Lys Phe Leu  
 130 135 140

gcc agc gtt acc cct gtc ggt gtc aat atg gcg aca gac atc gac aaa 2042  
 Ala Ser Val Thr Pro Val Gly Val Asn Met Ala Thr Asp Ile Asp Lys  
 145 150 155

tac ctc tcc ttc atc ttg ggg atg ttt gtc gca ttc ggt aca acg ttt 2090  
 Tyr Leu Ser Phe Ile Leu Gly Met Phe Val Ala Phe Gly Thr Thr Phe  
 160 165 170

gaa gtc ccc att gtc gtt atc ctg tta acc aaa att ggt gtg gta aca 2138  
 Glu Val Pro Ile Val Val Ile Leu Leu Thr Lys Ile Gly Val Val Thr  
 175 180 185

acc gaa cag ctc aaa cgc gcc cgc ccc tat gtg att gtc ggc gcg ttt 2186  
 Thr Glu Gln Leu Lys Arg Ala Arg Pro Tyr Val Ile Val Gly Ala Phe  
 190 195 200 205

gtc att gcc gcc atc atc acg ccg ccc gat gtg att tca caa acc ctg 2234  
 Val Ile Ala Ala Ile Ile Thr Pro Pro Asp Val Ile Ser Gln Thr Leu  
 210 215 220

ctt gcc att ccg ctg att ctc tta tac gaa gca ggt att tgg ttc gga 2282  
 Leu Ala Ile Pro Leu Ile Leu Leu Tyr Glu Ala Gly Ile Trp Phe Gly  
 225 230 235

cgc ttt ttc acg cca cgt tca gaa cag gat ggc gac ata cag ccg cct 2330  
 Arg Phe Phe Thr Pro Arg Ser Glu Gln Asp Gly Asp Ile Trp Phe Pro  
 240 245 250

gca aca acc tgacactatg ccgtccgaac ctccgcctca taccgccaca 2379  
 Ala Thr Thr  
 255

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<210> 21

<211> 256

<212> PRT

<213> Neisseria meningitidis

<400> 21

Val Ser Glu Thr Gln Asn Glu Gln Pro Val Gln Pro Leu Val Glu His  
 1 5 10 15

Leu Ile Glu Leu Arg Arg Arg Leu Met Trp Thr Val Val Gly Ile Leu  
 20 25 30

Val Cys Phe Phe Gly Leu Met Pro Phe Ala Gln Gln Leu Tyr Thr Phe  
 35 40 45

Ile Ala Asp Pro Leu Met Ala Asn Leu Pro Lys Asp Thr Ser Met Ile  
 50 55 60

Ala Thr Asp Val Ile Ala Pro Phe Phe Val Pro Val Lys Val Thr Leu  
 65 70 75 80

Met Ala Ala Phe Leu Ile Ser Leu Pro His Thr Leu Tyr Gln Ile Trp  
 85 90 95

Ala Phe Val Ala Pro Ala Leu Tyr Gln Asn Glu Lys Arg Leu Ile Thr  
 100 105 110

Pro Leu Val Leu Ser Ser Val Ser Leu Phe Phe Ile Gly Met Ala Phe  
 115 120 125

Ala Tyr Phe Leu Val Phe Pro Val Ile Phe Lys Phe Leu Ala Ser Val  
 130 135 140

Thr Pro Val Gly Val Asn Met Ala Thr Asp Ile Asp Lys Tyr Leu Ser  
 145 150 155 160



Phe Ile Leu Gly Met Phe Val Ala Phe Gly Thr Thr Phe Glu Val Pro  
 165 170 175

Ile Val Val Ile Leu Leu Thr Lys Ile Gly Val Val Thr Thr Glu Gln  
 180 185 190

Leu Lys Arg Ala Arg Pro Tyr Val Ile Val Gly Ala Phe Val Ile Ala  
 195 200 205

Ala Ile Ile Thr Pro Pro Asp Val Ile Ser Gln Thr Leu Leu Ala Ile  
 210 215 220

Pro Leu Ile Leu Leu Tyr Glu Ala Gly Ile Trp Phe Gly Arg Phe Phe  
 225 230 235 240

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<211> 4604

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 1 5 10 15

tcg ctg ata aag tac agc gag aca gat tat aca att tat tgt gac caa 98



Ser Phe Cys Glu Ser Asn Asn Lys Phe Thr Asp Phe Phe Lys Leu Trp	
210	215
cga ggt ggg ttt aga tta aat aac agt aga act aaa tta tta tta aaa	722
Arg Gly Gly Phe Arg Leu Asn Asn Ser Arg Thr Lys Leu Leu Lys	
225	230
ttc tta ata cgg aga aaa ttt agc ga atg att tca ata ctt aca cct	769
Phe Leu Ile Arg Arg Lys Phe Ser Met Ile Ser Ile Leu Thr Pro	
240	245
act ttt aat cgg caa cat act tta tca agg cta ttc aat tct ctt ata	817
Thr Phe Asn Arg Gln His Thr Leu Ser Arg Leu Phe Asn Ser Leu Ile	
255	260
tta caa act gat aaa gat ttt gag tgg ata ata att gat gat ggt agt	865
Leu Gln Thr Asp Lys Asp Phe Glu Trp Ile Ile Ile Asp Asp Gly Ser	
275	280
ata gat gca aca gcg gta ctt gta gaa gat ttt aga aaa aaa tgt gat	913
Ile Asp Ala Thr Ala Val Leu Val Glu Asp Phe Arg Lys Lys Cys Asp	
290	295
ttt gac ttg att tat tgc tat cag gaa aat aat ggt aag ccc atg gct	961
Phe Asp Leu Ile Tyr Cys Tyr Gln Glu Asn Asn Gly Lys Pro Met Ala	
305	310
tta aac gct ggt gtt aaa gct tgt aga ggc gat tat atc ttt att gtt	1009
Leu Asn Ala Gly Val Lys Ala Cys Arg Gly Asp Tyr Ile Phe Ile Val	
320	325
gac agt gat gat gca cta act ccc gat gcc ata aaa tta att aaa gaa	1057
Asp Ser Asp Asp Ala Leu Thr Pro Asp Ala Ile Lys Leu Ile Lys Glu	
335	340
tca ata cat gat tgc tta tct gag aag gaa agt ttc agc gga gtc ggt	1105
Ser Ile His Asp Cys Leu Ser Glu Lys Glu Ser Phe Ser Gly Val Gly	
355	360
ttt aga aaa gca tat ata aaa ggg ggg att att ggt aat gat tta aat	1153
Phe Arg Lys Ala Tyr Ile Lys Gly Gly Ile Ile Gly Asn Asp Leu Asn	
370	375
aat tct tca gaa cat ata tac tat tta aat gcg act gag att agc aat	1201
Asn Ser Ser Glu His Ile Tyr Tyr Leu Asn Ala Thr Glu Ile Ser Asn	
385	390
tta ata aat ggt gat gtt gca tat tgt ttt aaa aaa gaa agt ttg gta	1249

Leu Ile Asn Gly Asp Val Ala Tyr Cys Phe Lys Lys Glu Ser Leu Val	
400 405 410	
aaa aat cca ttc ccc cgt ata gaa gat gaa aaa ttt gtt cca gaa tta	1297
Lys Asn Pro Phe Pro Arg Ile Glu Asp Glu Lys Phe Val Pro Glu Leu	
415 420 425 430	
tat att tgg aat aaa ata act gac aag gcg aag att cga ttt aac ata	1345
Tyr Ile Trp Asn Lys Ile Thr Asp Lys Ala Lys Ile Arg Phe Asn Ile	
435 440 445	
agc aaa gtt ata tat ctt tgt gag tat ctt gat gat ggt ctt tct aaa	1393
Ser Lys Val Ile Tyr Leu Cys Glu Tyr Leu Asp Asp Gly Leu Ser Lys	
450 455 460	
aat ttc cat aac cag ctt aaa aaa tac cca aag ggg ttt aag att tat	1441
Asn Phe His Asn Gln Leu Lys Lys Tyr Pro Lys Gly Phe Lys Ile Tyr	
465 470 475	
tac aaa gat caa aga aaa cga gag aaa act tat ata aaa aaa aca aag	1489
Tyr Lys Asp Gln Arg Lys Arg Glu Lys Thr Tyr Ile Lys Lys Thr Lys	
480 485 490	
atg cta att aga tat ttg caa tgt tgt tat tat gag aaa ata aa atg	1536
Met Leu Ile Arg Tyr Leu Gln Cys Cys Tyr Tyr Glu Lys Ile Met	
495 500 505	
aaa ata cta ttt gtc att aca ggt tta ggc ctt gga ggt gct gag aag	1584
Lys Ile Leu Phe Val Ile Thr Gly Leu Gly Leu Gly Gly Ala Glu Lys	
510 515 520 525	
cag gtt tgt ctt tta gct gat aaa tta agt tta agc ggg cac cat gta	1632
Gln Val Cys Leu Leu Ala Asp Lys Leu Ser Leu Ser Gly His His Val	
530 535 540	
aag att att tca ctt gga cat atg tct aat aat aaa gtc ttt cct agc	1680
Lys Ile Ile Ser Leu Gly His Met Ser Asn Asn Lys Val Phe Pro Ser	
545 550 555	
gaa aat aat gtt aat gtc att aat gta aat atg tca aaa aac att tct	1728
Glu Asn Asn Val Asn Val Ile Asn Val Asn Met Ser Lys Asn Ile Ser	
560 565 570	
gga gtt ata aaa ggt tgt gtc aga att aga gat gtt ata gct aat ttc	1776
Gly Val Ile Lys Gly Cys Val Arg Ile Arg Asp Val Ile Ala Asn Phe	
575 580 585	
aaa cca gac att gta cac agt cat atg ttt cat gca aac att atc act	1824

Lys Pro Asp Ile Val His Ser His Met Phe His Ala Asn Ile Ile Thr	
590	595 600 605
aga ttg tct gta att gga atc aaa aac aga cct ggt att ata tca act	1872
Arg Leu Ser Val Ile Gly Ile Lys Asn Arg Pro Gly Ile Ile Ser Thr	
610 615 620	
gca cat aat aaa aat gaa ggt ggg tat ttc aga atg ctc aca tat aga	1920
Ala His Asn Lys Asn Glu Gly Gly Tyr Phe Arg Met Leu Thr Tyr Arg	
625 630 635	
ata acc gat tgt tta agt gat tgt tgt aca aat gtt agc aaa gaa gca	1968
Ile Thr Asp Cys Leu Ser Asp Cys Cys Thr Asn Val Ser Lys Glu Ala	
640 645 650	
gtg gat gag ttt tta cgg ata aaa gcc ttt aat ccc gct aaa gca att	2016
Val Asp Glu Phe Leu Arg Ile Lys Ala Phe Asn Pro Ala Lys Ala Ile	
655 660 665	
act atg tat aat ggg ata gat acc aat aaa ttt aaa ttt gat tta ttg	2064
Thr Met Tyr Asn Gly Ile Asp Thr Asn Lys Phe Lys Phe Asp Leu Leu	
670 675 680 685	
gca agg agg gaa att cga gac ggt att aat ata aaa aat gat gat ata	2112
Ala Arg Arg Glu Ile Arg Asp Gly Ile Asn Ile Lys Asn Asp Asp Ile	
690 695 700	
tta tta ctt gct gca ggt cgt tta acg tta gct aaa gat tat cct aat	2160
Leu Leu Leu Ala Ala Gly Arg Leu Thr Leu Ala Lys Asp Tyr Pro Asn	
705 710 715	
tta ttg aat gca atg act ctg ctt cct gaa cac ttt aaa ctt att att	2208
Leu Leu Asn Ala Met Thr Leu Leu Pro Glu His Phe Lys Leu Ile Ile	
720 725 730	
att ggt gat ggt gaa ttg cgt gac gaa att aat atg ctt ata aaa aaa	2256
Ile Gly Asp Gly Glu Leu Arg Asp Glu Ile Asn Met Leu Ile Lys Lys	
735 740 745	
ttg caa tta tct aat agg gtg tcc ttg ttg gga gtt aaa aaa aat att	2304
Leu Gln Leu Ser Asn Arg Val Ser Leu Leu Gly Val Lys Lys Asn Ile	
750 755 760 765	
gct ccc tat ttt tct gca tgt gat att ttt gtt ctc tct tct cgt tgg	2352
Ala Pro Tyr Phe Ser Ala Cys Asp Ile Phe Val Leu Ser Ser Arg Trp	
770 775 780	
gaa gga ttt gga tta gtc gtg gca gaa gct atg tca tgt gag cga att	2400

Glu Gly Phe Gly Leu Val Val Ala	Glu Ala Met Ser Cys Glu Arg Ile	
785	790	795
gtt gtt ggc acg gat tca ggg gga gta aga gaa gtt att ggt gac gat	2448	
Val Val Gly Thr Asp Ser Gly Gly Val Arg Glu Val Ile Gly Asp Asp		
800	805	810
gat ttt ctt gta ccc ata tct gat tca aca caa ctt gca agc aaa att	2496	
Asp Phe Leu Val Pro Ile Ser Asp Ser Thr Gln Leu Ala Ser Lys Ile		
815	820	825
gaa aaa ttg tct ttg agc cag ata cgt gat cac att ggt ttt cgg aat	2544	
Glu Lys Leu Ser Leu Ser Gln Ile Arg Asp His Ile Gly Phe Arg Asn		
830	835	840 845
cgt gag cgt att tta aaa aat ttc tca ata gat act att att atg cag	2592	
Arg Glu Arg Ile Leu Lys Asn Phe Ser Ile Asp Thr Ile Ile Met Gln		
850	855	860
tg g caa gaa ct c tat gga act ata att tgc tca aaa cat gaa agg	2637	
Trp Gln Glu Leu Tyr Gly Thr Ile Ile Cys Ser Lys His Glu Arg		
865	870	875
tagatttata tttggaacgt gtcttttgtt tgaatttaaat tcaatctcaa ttgagatttt	2697	
tgtatttcaa aaataccatc atagctaacg atgattggta tttatttttaa gatgctttct	2757	
ataaatatat tgacgttttt aatgcgccga aacgattggg ctgggaacag agaagtaaaa	2817	
ctgttttgag aatgaagagt ttttgagatg tttatggata ttaaaaattg atccagttaa	2877	
ttaattattt ataataaatc aagattttaat gttaataaat gataatcttt tctgacactc	2937	
atattaatta tgagtgttac gtttggtaaa cggtaaaacta ttat atg aca gct aga	2993	
	Met Thr Ala Arg	
	880	
aca act aaa gtt ttg cac tta caa tta ct c cca ct c tta agt ggc gtt	3041	
Thr Thr Lys Val Leu His Leu Gln Leu Leu Pro Leu Leu Ser Gly Val		
885	890	895
caa agg gta aca tta aac gaa att agt gcg tta tat act gat tat gat	3089	
Gln Arg Val Thr Leu Asn Glu Ile Ser Ala Leu Tyr Thr Asp Tyr Asp		
900	905	910
tat aca cta gtt tgc tca aaa aaa ggt cca cta aca aaa gca ttg ctg	3137	
Tyr Thr Leu Val Cys Ser Lys Lys Gly Pro Leu Thr Lys Ala Leu Leu		
915	920	925

gaa tat gat gtc gat tgt cat tgt atc ccc gaa ctt acg aga gaa att	3185
Glu Tyr Asp Val Asp Cys His Cys Ile Pro Glu Leu Thr Arg Glu Ile	
930 935 940	
acc gta aag aat gat ttt aaa gca ttg ttc aag ctt tat aag ttc ata	3233
Thr Val Lys Asn Asp Phe Lys Ala Leu Phe Lys Leu Tyr Lys Phe Ile	
945 950 955 960	
aaa aaa gaa aaa ttt gac att gtg cat aca cat tct tca aaa aca ggt	3281
Lys Lys Glu Lys Phe Asp Ile Val His Thr His Ser Ser Lys Thr Gly	
965 970 975	
att ttg ggg cga gtt gct gcc aaa tta gca cgt gtt gga aag gtg atc	3329
Ile Leu Gly Arg Val Ala Ala Lys Leu Ala Arg Val Gly Lys Val Ile	
980 985 990	
cac act gta cat ggt ttt tct ttt cca gcc gca tct agt aaa aaa agt	3377
His Thr Val His Gly Phe Ser Phe Pro Ala Ala Ser Ser Lys Lys Ser	
995 1000 1005	
tat tac ctt tat ttt ttc atg gaa tgg ata gca aag ttc ttt acg gat	3425
Tyr Tyr Leu Tyr Phe Phe Met Glu Trp Ile Ala Lys Phe Phe Thr Asp	
1010 1015 1020	
aag tta atc gtc ttg aat gta gat gat gaa tat ata gca ata aac aaa	3473
Lys Leu Ile Val Leu Asn Val Asp Asp Glu Tyr Ile Ala Ile Asn Lys	
1025 1030 1035 1040	
tta aaa ttc aag cgg gat aaa gtt ttt tta att cct aat gga gta gac	3521
Leu Lys Phe Lys Arg Asp Lys Val Phe Leu Ile Pro Asn Gly Val Asp	
1045 1050 1055	
act gat aag ttt tct cct tta gaa aat aaa att tat agt agc acc ttg	3569
Thr Asp Lys Phe Ser Pro Leu Glu Asn Lys Ile Tyr Ser Ser Thr Leu	
1060 1065 1070	
aat cta gta atg gtt ggt aga tta tcc aag caa aaa gat cct gag aca	3617
Asn Leu Val Met Val Gly Arg Leu Ser Lys Gln Lys Asp Pro Glu Thr	
1075 1080 1085	
tta ttg ctt gct gtt gaa aaa ctg ctg aat gaa aat gtt aat gtt aag	3665
Leu Leu Leu Ala Val Glu Lys Leu Leu Asn Glu Asn Val Asn Val Lys	
1090 1095 1100	
ctg aca ctt gta gga gat ggt gaa cta aaa gaa cag tta gaa agc agg	3713
Leu Thr Leu Val Gly Asp Gly Glu Leu Lys Glu Gln Leu Glu Ser Arg	
1105 1110 1115 1120	

ttc aaa cgg caa gat gga cgt ata att ttt cat gga tgg tca gat aac 3761  
 Phe Lys Arg Gln Asp Gly Arg Ile Ile Phe His Gly Trp Ser Asp Asn  
 1125 1130 1135

att gtt aat att tta aaa gtt aat gat ctt ttt ata tta cct tct ctt 3809  
 Ile Val Asn Ile Leu Lys Val Asn Asp Leu Phe Ile Leu Pro Ser Leu  
 1140 1145 1150

tgg gag ggt atg cca tta gca att tta gaa gca ttg agc tgt gga ctt 3857  
 Trp Glu Gly Met Pro Leu Ala Ile Leu Glu Ala Leu Ser Cys Gly Leu  
 1155 1160 1165

cca tgt ata gtc act aat att cca ggt aat aat agc tta ata gaa gat 3905  
 Pro Cys Ile Val Thr Asn Ile Pro Gly Asn Asn Ser Leu Ile Glu Asp  
 1170 1175 1180

ggc tat aat ggt tgt ttg ttt gaa att aga gat tgt cag tta tta tct 3953  
 Gly Tyr Asn Gly Cys Leu Phe Glu Ile Arg Asp Cys Gln Leu Leu Ser  
 1185 1190 1195 1200

caa aaa atc atg tca tat gtt ggt aag cca gaa ctg att gca cag caa 4001  
 Gln Lys Ile Met Ser Tyr Val Gly Lys Pro Glu Leu Ile Ala Gln Gln  
 1205 1210 1215

tct acc aat gca cga tca ttt att ctg aaa aat tat gga tta gtt aaa 4049  
 Ser Thr Asn Ala Arg Ser Phe Ile Leu Lys Asn Tyr Gly Leu Val Lys  
 1220 1225 1230

aga aat aat aag gtc aga cag cta tat gat aat taaatgaaac cgaaaagtta 4102  
 Arg Asn Asn Lys Val Arg Gln Leu Tyr Asp Asn  
 1235 1240

aaaaagaaca ggtttttcaa agtgaaaata aaattacagt ttttttattg caatgattaa 4162

cgtaacatct gcattacatt caagcgcgac aaccccgcg tgaccacccc tgacaggagt 4222

aaacaatgtc aaagcaacag atcggcgtcg tcggtatggc agtgatggga cgcaacctcg 4282

cgctcaacat cgaaagccgt ggttataccg tctctatatt caaccgttcc cgtgaaaaga 4342

cggaagaagt tattgccgaa aatccaggca agaaactggt tccttactat acggtgaaag 4402

agttcgttga atctcttgaa acgcctcgtc gcatcctggt aatgggttaa agcaggtgca 4462

ggcaaggatg ctgctattga ttccctgaaa ccatatctcg ataaaggcga tatcatcatt 4522

gatgggtggg taataccttc ttccaggaca ccattcgtcg taaccgcgag ctttctgcac 4582



aaggctttac ttcacggta cc

4604

&lt;210&gt; 23

&lt;211&gt; 247

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 23

Val Asp Asp Asn Leu Lys Gly Gln Gly Ala Gly Lys Asn Phe Leu Ser  
 1 5 10 15

Leu Ile Lys Tyr Ser Glu Thr Asp Tyr Thr Ile Tyr Cys Asp Gln Asp  
 20 25 30

Asp Ile Trp Leu Glu Asn Lys Ile Phe Glu Leu Val Lys Tyr Ala Asn  
 35 40 45

Glu Ile Lys Leu Asn Val Ser Asp Ala Pro Ser Leu Val Tyr Ala Asp  
 50 55 60

Gly Tyr Ala Tyr Met Asp Gly Glu Gly Thr Ile Asp Phe Ser Gly Ile  
 65 70 75 80

Ser Asn Asn His Ala Asp Gln Leu Lys Asp Phe Leu Phe Phe Asn Gly  
 85 90 95

Gly Tyr Gln Gly Cys Ser Ile Met Phe Asn Arg Ala Met Thr Lys Phe  
 100 105 110

Leu Leu Asn Tyr Arg Gly Phe Val Tyr Leu His Asp Asp Ile Thr Thr  
 115 120 125

Leu Ala Ala Tyr Ala Leu Gly Lys Val Tyr Phe Leu Pro Lys Tyr Leu  
 130 135 140

Met Leu Tyr Arg Gln His Thr Asn Ala Val Thr Gly Ile Lys Thr Phe  
 145 150 155 160

Arg Asn Gly Leu Thr Ser Lys Phe Lys Ser Pro Val Asn Tyr Leu Leu  
 165 170 175

Ser Arg Lys His Tyr Gln Val Lys Lys Ser Phe Phe Glu Cys Asn Ser  
 180 185 190

Ser Ile Leu Ser Glu Thr Asn Lys Lys Val Phe Leu Asp Phe Ile Ser  
 195 200 205

Phe Cys Glu Ser Asn Asn Lys Phe Thr Asp Phe Phe Lys Leu Trp Arg  
 210 215 220

Gly Gly Phe Arg Leu Asn Asn Ser Arg Thr Lys Leu Leu Leu Lys Phe  
 225 230 235 240

Leu Ile Arg Arg Lys Phe Ser  
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<211> 261

<212> PRT

<213> Escherichia coli

<400> 24

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Arg Leu Phe Asn Ser Leu Ile Leu Gln Thr Asp Lys Asp Phe Glu Trp  
 20 25 30

Ile Ile Ile Asp Asp Gly Ser Ile Asp Ala Thr Ala Val Leu Val Glu  
 35 40 45

Asp Phe Arg Lys Lys Cys Asp Phe Asp Leu Ile Tyr Cys Tyr Gln Glu  
 50 55 60

Asn Asn Gly Lys Pro Met Ala Leu Asn Ala Gly Val Lys Ala Cys Arg  
 65 70 75 80

Gly Asp Tyr Ile Phe Ile Val Asp Ser Asp Asp Ala Leu Thr Pro Asp  
 85 90 95

Ala Ile Lys Leu Ile Lys Glu Ser Ile His Asp Cys Leu Ser Glu Lys  
 100 105 110

Glu Ser Phe Ser Gly Val Gly Phe Arg Lys Ala Tyr Ile Lys Gly Gly  
 115 120 125

Ile Ile Gly Asn Asp Leu Asn Asn Ser Ser Glu His Ile Tyr Tyr Leu  
 130 135 140

Asn Ala Thr Glu Ile Ser Asn Leu Ile Asn Gly Asp Val Ala Tyr Cys  
 145 150 155 160

Phe Lys Lys Glu Ser Leu Val Lys Asn Pro Phe Pro Arg Ile Glu Asp

165                      170                      175

Glu Lys Phe Val Pro Glu Leu Tyr Ile Trp Asn Lys Ile Thr Asp Lys  
180                      185                      190

Ala Lys Ile Arg Phe Asn Ile Ser Lys Val Ile Tyr Leu Cys Glu Tyr  
195                      200                      205

Leu Asp Asp Gly Leu Ser Lys Asn Phe His Asn Gln Leu Lys Lys Tyr  
210                      215                      220

Pro Lys Gly Phe Lys Ile Tyr Tyr Lys Asp Gln Arg Lys Arg Glu Lys  
225                      230                      235                      240

Thr Tyr Ile Lys Lys Thr Lys Met Leu Ile Arg Tyr Leu Gln Cys Cys  
245                      250                      255

Tyr Tyr Glu Lys Ile  
260

<210> 25  
<211> 368  
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<400> 25

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Lys Gln Val Cys Leu Leu Ala Asp Lys Leu Ser Leu Ser Gly His His  
20                      25                      30

Val Lys Ile Ile Ser Leu Gly His Met Ser Asn Asn Lys Val Phe Pro  
35                      40                      45

Ser Glu Asn Asn Val Asn Val Ile Asn Val Asn Met Ser Lys Asn Ile  
50                      55                      60

Ser Gly Val Ile Lys Gly Cys Val Arg Ile Arg Asp Val Ile Ala Asn  
65                      70                      75                      80

Phe Lys Pro Asp Ile Val His Ser His Met Phe His Ala Asn Ile Ile  
85                      90                      95

Thr Arg Leu Ser Val Ile Gly Ile Lys Asn Arg Pro Gly Ile Ile Ser  
100                      105                      110

Thr Ala His Asn Lys Asn Glu Gly Gly Tyr Phe Arg Met Leu Thr Tyr  
 115 120 125

Arg Ile Thr Asp Cys Leu Ser Asp Cys Cys Thr Asn Val Ser Lys Glu  
 130 135 140

Ala Val Asp Glu Phe Leu Arg Ile Lys Ala Phe Asn Pro Ala Lys Ala  
 145 150 155 160

Ile Thr Met Tyr Asn Gly Ile Asp Thr Asn Lys Phe Lys Phe Asp Leu  
 165 170 175

Leu Ala Arg Arg Glu Ile Arg Asp Gly Ile Asn Ile Lys Asn Asp Asp  
 180 185 190

Ile Leu Leu Leu Ala Ala Gly Arg Leu Thr Leu Ala Lys Asp Tyr Pro  
 195 200 205

Asn Leu Leu Asn Ala Met Thr Leu Leu Pro Glu His Phe Lys Leu Ile  
 210 215 220

Ile Ile Gly Asp Gly Glu Leu Arg Asp Glu Ile Asn Met Leu Ile Lys  
 225 230 235 240

Lys Leu Gln Leu Ser Asn Arg Val Ser Leu Leu Gly Val Lys Lys Asn  
 245 250 255

Ile Ala Pro Tyr Phe Ser Ala Cys Asp Ile Phe Val Leu Ser Ser Arg  
 260 265 270

Trp Glu Gly Phe Gly Leu Val Val Ala Glu Ala Met Ser Cys Glu Arg  
 275 280 285

Ile Val Val Gly Thr Asp Ser Gly Gly Val Arg Glu Val Ile Gly Asp  
 290 295 300

Asp Asp Phe Leu Val Pro Ile Ser Asp Ser Thr Gln Leu Ala Ser Lys  
 305 310 315 320

Ile Glu Lys Leu Ser Leu Ser Gln Ile Arg Asp His Ile Gly Phe Arg  
 325 330 335

Asn Arg Glu Arg Ile Leu Lys Asn Phe Ser Ile Asp Thr Ile Ile Met  
 340 345 350

Gln Trp Gln Glu Leu Tyr Gly Thr Ile Ile Cys Ser Lys His Glu Arg  
 355 360 365

&lt;210&gt; 26

&lt;211&gt; 367

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 26

Met Thr Ala Arg Thr Thr Lys Val Leu His Leu Gln Leu Leu Pro Leu  
 1 5 10 15

Leu Ser Gly Val Gln Arg Val Thr Leu Asn Glu Ile Ser Ala Leu Tyr  
 20 25 30

Thr Asp Tyr Asp Tyr Thr Leu Val Cys Ser Lys Lys Gly Pro Leu Thr  
 35 40 45

Lys Ala Leu Leu Glu Tyr Asp Val Asp Cys His Cys Ile Pro Glu Leu  
 50 55 60

Thr Arg Glu Ile Thr Val Lys Asn Asp Phe Lys Ala Leu Phe Lys Leu  
 65 70 75 80

Tyr Lys Phe Ile Lys Lys Glu Lys Phe Asp Ile Val His Thr His Ser  
 85 90 95

Ser Lys Thr Gly Ile Leu Gly Arg Val Ala Ala Lys Leu Ala Arg Val  
 100 105 110

Gly Lys Val Ile His Thr Val His Gly Phe Ser Phe Pro Ala Ala Ser  
 115 120 125

Ser Lys Lys Ser Tyr Tyr Leu Tyr Phe Phe Met Glu Trp Ile Ala Lys  
 130 135 140

Phe Phe Thr Asp Lys Leu Ile Val Leu Asn Val Asp Asp Glu Tyr Ile  
 145 150 155 160

Ala Ile Asn Lys Leu Lys Phe Lys Arg Asp Lys Val Phe Leu Ile Pro  
 165 170 175

Asn Gly Val Asp Thr Asp Lys Phe Ser Pro Leu Glu Asn Lys Ile Tyr  
 180 185 190

Ser Ser Thr Leu Asn Leu Val Met Val Gly Arg Leu Ser Lys Gln Lys  
 195 200 205

Asp Pro Glu Thr Leu Leu Leu Ala Val Glu Lys Leu Leu Asn Glu Asn  
 210 215 220

Val Asn Val Lys Leu Thr Leu Val Gly Asp Gly Glu Leu Lys Glu Gln  
 225 230 235 240

Leu Glu Ser Arg Phe Lys Arg Gln Asp Gly Arg Ile Ile Phe His Gly  
 245 250 255

Trp Ser Asp Asn Ile Val Asn Ile Leu Lys Val Asn Asp Leu Phe Ile  
 260 265 270

Leu Pro Ser Leu Trp Glu Gly Met Pro Leu Ala Ile Leu Glu Ala Leu  
 275 280 285

Ser Cys Gly Leu Pro Cys Ile Val Thr Asn Ile Pro Gly Asn Asn Ser  
 290 295 300

Leu Ile Glu Asp Gly Tyr Asn Gly Cys Leu Phe Glu Ile Arg Asp Cys  
 305 310 315 320

Gln Leu Leu Ser Gln Lys Ile Met Ser Tyr Val Gly Lys Pro Glu Leu  
 325 330 335

Ile Ala Gln Gln Ser Thr Asn Ala Arg Ser Phe Ile Leu Lys Asn Tyr  
 340 345 350

Gly Leu Val Lys Arg Asn Asn Lys Val Arg Gln Leu Tyr Asp Asn  
 355 360 365

<210> 27

<211> 1272

<212> DNA

<213> Escherichia coli

<220>

<221> CDS

<222> (319)..(1269)

<220>

<221> CDS

<222> (3)..(215)

<400> 27

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 1 5 10 15

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caa cgc tgc gat ctg att gcc gtt att gat aag ggg tta ctt gcg gaa   95
Gln Arg Cys Asp Leu Ile Ala Val Ile Asp Lys Gly Leu Leu Ala Glu
                20                25                30

tac gga acc cac gaa cag ctg tta tct gcg ggc ggc ctc tat acc cgc   143
Tyr Gly Thr His Glu Gln Leu Leu Ser Ala Gly Gly Leu Tyr Thr Arg
                35                40                45

tta tgg cat gac agc gtc agc agt act gct ctc cat cgc cag cac aac   191
Leu Trp His Asp Ser Val Ser Ser Thr Ala Leu His Arg Gln His Asn
                50                55                60

atg aag gag gaa acc ccg gga tag ttactggaca cgtaatgtat taaaacaca   245
Met Lys Glu Glu Thr Pro Gly
                65                70

gtcagaagcgc gcggtaccgt gaatagccgc ttttaattatt tatactgaca tccttaattt 305

ttaaagagta tga atg ctg aac atg caa caa cat ctc tct gct atc gcc   354
                Met Leu Asn Met Gln Gln His Leu Ser Ala Ile Ala
                        75                        80

agc ctg cgc aac caa ctg gca gcg ggc cac att gct aac ctt act gac   402
Ser Leu Arg Asn Gln Leu Ala Ala Gly His Ile Ala Asn Leu Thr Asp
                85                90                95

ttc tgg cgc gaa gct gag tcg ctg aat gtt cct ctt gtg acg cca gtc   450
Phe Trp Arg Glu Ala Glu Ser Leu Asn Val Pro Leu Val Thr Pro Val
100                105                110                115

gaa gga gcg gaa gat gag cga gaa gtg acc ttt ctg tgg cgc gcc cga   498
Glu Gly Ala Glu Asp Glu Arg Glu Val Thr Phe Leu Trp Arg Ala Arg
                120                125                130

cat cct ctg cag ggc gtt tat ctg cgt ctg aac cgg gtg acg gat aaa   546
His Pro Leu Gln Gly Val Tyr Leu Arg Leu Asn Arg Val Thr Asp Lys
                135                140                145

gag cac gta gaa aaa gga atg atg agc gcc ctt ccc gaa acg gat atc   594
Glu His Val Glu Lys Gly Met Met Ser Ala Leu Pro Glu Thr Asp Ile
                150                155                160

tgg aca ctg aca ctg cgt tta ccc gca agt tac tgc ggc tcc tat tcg   642
Trp Thr Leu Thr Leu Arg Leu Pro Ala Ser Tyr Cys Gly Ser Tyr Ser
                165                170                175

ctg ctg gaa atc ccc ccc ggc act acg gct gag acg att gca ctg tcc   690
Leu Leu Glu Ile Pro Pro Gly Thr Thr Ala Glu Thr Ile Ala Leu Ser

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180	185	190	195	
gga ggc cgt ttt gcc acc ctt gcc gga aag gcc gat ccg cta aac aaa				738
Gly Gly Arg Phe Ala Thr Leu Ala Gly Lys Ala Asp Pro Leu Asn Lys				
200		205	210	
atg ccg gag atc aac gtt cgg gga aac gca aag gaa tca gtg ctg aca				786
Met Pro Glu Ile Asn Val Arg Gly Asn Ala Lys Glu Ser Val Leu Thr				
215	220		225	
ctt gat aaa gct ccc gcc ctg tcg gaa tgg aac ggc ggc ttc cac acc				834
Leu Asp Lys Ala Pro Ala Leu Ser Glu Trp Asn Gly Gly Phe His Thr				
230	235		240	
gga caa ctg ctt acc tcc atg cgc att atc gcc ggg aaa tct cgc cag				882
Gly Gln Leu Leu Thr Ser Met Arg Ile Ile Ala Gly Lys Ser Arg Gln				
245	250		255	
gtt cgg ctc tat att ccg gat gtt gat att tct cag ccc ctc ggg ctg				930
Val Arg Leu Tyr Ile Pro Asp Val Asp Ile Ser Gln Pro Leu Gly Leu				
260	265	270	275	
gtc gtg ctg ccc gat ggt gaa acc tgg ttt gat cac ctt ggc gta tgc				978
Val Val Leu Pro Asp Gly Glu Thr Trp Phe Asp His Leu Gly Val Cys				
280	285		290	
gcg gca att gac gcc gcc ata aat aat ggg cgc atc gtg ccc gtg gct				1026
Ala Ala Ile Asp Ala Ala Ile Asn Asn Gly Arg Ile Val Pro Val Ala				
295	300		305	
gta ctg ggc att gac aac att aat gaa cat gaa cgc act gag ata ctc				1074
Val Leu Gly Ile Asp Asn Ile Asn Glu His Glu Arg Thr Glu Ile Leu				
310	315		320	
ggc ggg cgc agc aaa ctg ata aag gat atc gcc gga cat ctg ctg ccg				1122
Gly Gly Arg Ser Lys Leu Ile Lys Asp Ile Ala Gly His Leu Leu Pro				
325	330		335	
atg att cgc gct gaa caa ccg cag cgt cag tgg gca gac cgt tcg cgc				1170
Met Ile Arg Ala Glu Gln Pro Gln Arg Gln Trp Ala Asp Arg Ser Arg				
340	345	350	355	
aca gtg ctg gcc ggg cag agc ctc ggc ggg atc agt gcg cta atg ggg				1218
Thr Val Leu Ala Gly Gln Ser Leu Gly Gly Ile Ser Ala Leu Met Gly				
360	365		370	
gct cgt tac gca ccg gaa acg ttc ggt ctg gtg ctc agc cac tct cct				1266
Ala Arg Tyr Ala Pro Glu Thr Phe Gly Leu Val Leu Ser His Ser Pro				



375

380

385

caa tgc

1272

Gln

&lt;210&gt; 28

&lt;211&gt; 70

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 28

Gly Lys His Ser Ala Leu Ile Val Ala His Arg Leu Thr Thr Ala Gln

1

5

10

15

Arg Cys Asp Leu Ile Ala Val Ile Asp Lys Gly Leu Leu Ala Glu Tyr

20

25

30

Gly Thr His Glu Gln Leu Leu Ser Ala Gly Gly Leu Tyr Thr Arg Leu

35

40

45

Trp His Asp Ser Val Ser Ser Thr Ala Leu His Arg Gln His Asn Met

50

55

60

Lys Glu Glu Thr Pro Gly

65

70

&lt;210&gt; 29

&lt;211&gt; 317

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 29

Met Leu Asn Met Gln Gln His Leu Ser Ala Ile Ala Ser Leu Arg Asn

1

5

10

15

Gln Leu Ala Ala Gly His Ile Ala Asn Leu Thr Asp Phe Trp Arg Glu

20

25

30

Ala Glu Ser Leu Asn Val Pro Leu Val Thr Pro Val Glu Gly Ala Glu

35

40

45

Asp Glu Arg Glu Val Thr Phe Leu Trp Arg Ala Arg His Pro Leu Gln

50

55

60

Gly Val Tyr Leu Arg Leu Asn Arg Val Thr Asp Lys Glu His Val Glu

65

70

75

80

Lys Gly Met Met Ser Ala Leu Pro Glu Thr Asp Ile Trp Thr Leu Thr  
                                   85                                  90                                  95

Leu Arg Leu Pro Ala Ser Tyr Cys Gly Ser Tyr Ser Leu Leu Glu Ile  
                                   100                                  105                                  110

Pro Pro Gly Thr Thr Ala Glu Thr Ile Ala Leu Ser Gly Gly Arg Phe  
                                   115                                  120                                  125

Ala Thr Leu Ala Gly Lys Ala Asp Pro Leu Asn Lys Met Pro Glu Ile  
                                   130                                  135                                  140

Asn Val Arg Gly Asn Ala Lys Glu Ser Val Leu Thr Leu Asp Lys Ala  
                                   145                                  150                                  155                                  160

Pro Ala Leu Ser Glu Trp Asn Gly Gly Phe His Thr Gly Gln Leu Leu  
                                   165                                  170                                  175

Thr Ser Met Arg Ile Ile Ala Gly Lys Ser Arg Gln Val Arg Leu Tyr  
                                   180                                  185                                  190

Ile Pro Asp Val Asp Ile Ser Gln Pro Leu Gly Leu Val Val Leu Pro  
                                   195                                  200                                  205

Asp Gly Glu Thr Trp Phe Asp His Leu Gly Val Cys Ala Ala Ile Asp  
                                   210                                  215                                  220

Ala Ala Ile Asn Asn Gly Arg Ile Val Pro Val Ala Val Leu Gly Ile  
                                   225                                  230                                  235                                  240

Asp Asn Ile Asn Glu His Glu Arg Thr Glu Ile Leu Gly Gly Arg Ser  
                                   245                                  250                                  255

Lys Leu Ile Lys Asp Ile Ala Gly His Leu Leu Pro Met Ile Arg Ala  
                                   260                                  265                                  270

Glu Gln Pro Gln Arg Gln Trp Ala Asp Arg Ser Arg Thr Val Leu Ala  
                                   275                                  280                                  285

Gly Gln Ser Leu Gly Gly Ile Ser Ala Leu Met Gly Ala Arg Tyr Ala  
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Pro Glu Thr Phe Gly Leu Val Leu Ser His Ser Pro Gln  
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agc gaa acc gat acc tca tgg gtg agt gag cat ctg ctt tct gcc cca 96  
 Ser Glu Thr Asp Thr Ser Trp Val Ser Glu His Leu Leu Ser Ala Pro  
 20 25 30

ccg cag ggc gta cgt atc agc ctg tgc gtg gga tgc ctg gaa ggt tgc 144  
 Pro Gln Gly Val Arg Ile Ser Leu Cys Val Gly Ser Leu Glu Gly Ser  
 35 40 45

aca gtg cct cac gtt cag cag ctt cac cag cgg ctg att acc gct ggc 192  
 Thr Val Pro His Val Gln Gln Leu His Gln Arg Leu Ile Thr Ala Gly  
 50 55 60

gtc gaa agc cat tgc gca atc tac acc ggt ggt cac gat tac gca tgg 240  
 Val Glu Ser His Cys Ala Ile Tyr Thr Gly Gly His Asp Tyr Ala Trp  
 65 70 75 80

tgg cgc ggt gca ctg att gac ggg att ggt tta cta cag ggt tga 285  
 Trp Arg Gly Ala Leu Ile Asp Gly Ile Gly Leu Leu Gln Gly  
 85 90 95

gttgacccac aaacactttc aggaacggt acagacttcc tgaataaatc aaatagtcac 345

ctgcggaataa ggaataatca tcag atg tat gcc cgc gag tat cgc tca aca 396  
 Met Tyr Ala Arg Glu Tyr Arg Ser Thr  
 100

cgc ccg cat aaa gcg att ttc ttt cat ctt tct tgc ctc acc ctt atc 444  
 Arg Pro His Lys Ala Ile Phe Phe His Leu Ser Cys Leu Thr Leu Ile  
 105 110 115 120

tgt agt gcg caa gtt tat gcg aag ccg gat atg ccg cca ctg ggg ccg	492
Cys Ser Ala Gln Val Tyr Ala Lys Pro Asp Met Arg Pro Leu Gly Pro	
125 130 135	
aat ata gcc gat aaa ggc tcc gtg ttt tac cat ttc agc gtc acc tct	540
Asn Ile Ala Asp Lys Gly Ser Val Phe Tyr His Phe Ser Val Thr Ser	
140 145 150	
ttc gac tct gtc gat ggc aca cgc cat tat ccg gta tgg acg gcc gtg	588
Phe Asp Ser Val Asp Gly Thr Arg His Tyr Arg Val Trp Thr Ala Val	
155 160 165	
ccg aat aca acc gca ccg gca tcg ggt tac ccg att tta tat atg ctt	636
Pro Asn Thr Thr Ala Pro Ala Ser Gly Tyr Pro Ile Leu Tyr Met Leu	
170 175 180	
gac ggt aac gca gtt atg gat cgc ctg gat gac gaa ctg ctc aaa caa	684
Asp Gly Asn Ala Val Met Asp Arg Leu Asp Asp Glu Leu Leu Lys Gln	
185 190 195 200	
ttg tca gaa aaa aca ccg cca gtg atc gtg gct gtc ggg tat cag acc	732
Leu Ser Glu Lys Thr Pro Pro Val Ile Val Ala Val Gly Tyr Gln Thr	
205 210 215	
aac ctc cct ttc gat ctc aac agc agg gct tac gac tat acg cca gca	780
Asn Leu Pro Phe Asp Leu Asn Ser Arg Ala Tyr Asp Tyr Thr Pro Ala	
220 225 230	
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Ala Glu Ser Arg Lys Thr Asp Leu His Ser Gly Arg Phe Ser Arg Lys	
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agt ggt ggc agc aac aac ttc cgc cag tta ctg gaa acg cgt att gcc	876
Ser Gly Gly Ser Asn Asn Phe Arg Gln Leu Leu Glu Thr Arg Ile Ala	
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Pro Lys Val Glu Gln Gly Leu Asn Ile Asp Arg Gln Arg Arg Gly Leu	
265 270 275 280	
tgg ggg cac tcc tac ggc ggc ctc ttc gtg ctg gat tcc tgg ctg tcc	972
Trp Gly His Ser Tyr Gly Gly Leu Phe Val Leu Asp Ser Trp Leu Ser	
285 290 295	
tcc tct tac ttc ccg tcg tac tac agc gcc agc ccg tcg ttg ggc aga	1020
Ser Ser Tyr Phe Arg Ser Tyr Tyr Ser Leu Ser Pro Ser Leu Gly Arg	
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ggt tat gat gct ttg cta agc cgc gtt acg gcg gtt gag cct ctg caa 1068  
 Gly Tyr Asp Ala Leu Leu Ser Arg Val Thr Ala Val Glu Pro Leu Gln  
 315 320 325

ttc tgc gcc aaa cac ctg gcg ata atg gaa gcc tcg gcg aca cag ggt 1116  
 Phe Cys Ala Lys His Leu Ala Ile Met Glu Gly Ser Ala Thr Gln Gly  
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gat aac cgg gaa acg cat gct gtc ggg gtg ctg tcg aaa att cat acc 1164  
 Asp Asn Arg Glu Thr His Ala Val Gly Val Leu Ser Lys Ile His Thr  
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acc ctc act ata ctg aaa gat aaa ggc gtc aat gcc gta ttt tgg gat 1212  
 Thr Leu Thr Ile Leu Lys Asp Lys Gly Val Asn Ala Val Phe Trp Asp  
 365 370 375

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 Phe Pro Asn Leu Gly His Gly Pro Met Phe Asn Ala Ser Phe Arg Gln  
 380 385 390

gca ctg tta gat atc agt ggt gaa aac gca aat tac aca gca ggt tgt 1308  
 Ala Leu Leu Asp Ile Ser Gly Glu Asn Ala Asn Tyr Thr Ala Gly Cys  
 395 400 405

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&lt;210&gt; 31

&lt;211&gt; 94

&lt;212&gt; PRT

<213> *Escherichia coli*

&lt;400&gt; 31

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Ser Glu Thr Asp Thr Ser Trp Val Ser Glu His Leu Leu Ser Ala Pro  
 20 25 30

Pro Gln Gly Val Arg Ile Ser Leu Cys Val Gly Ser Leu Glu Gly Ser  
 35 40 45

Thr Val Pro His Val Gln Gln Leu His Gln Arg Leu Ile Thr Ala Gly  
 50 55 60

Val Glu Ser His Cys Ala Ile Tyr Thr Gly Gly His Asp Tyr Ala Trp  
 65 70 75 80

Trp Arg Gly Ala Leu Ile Asp Gly Ile Gly Leu Leu Gln Gly  
 85 90

&lt;210&gt; 32

&lt;211&gt; 318

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 32

Met Tyr Ala Arg Glu Tyr Arg Ser Thr Arg Pro His Lys Ala Ile Phe  
 1 5 10 15

Phe His Leu Ser Cys Leu Thr Leu Ile Cys Ser Ala Gln Val Tyr Ala  
 20 25 30

Lys Pro Asp Met Arg Pro Leu Gly Pro Asn Ile Ala Asp Lys Gly Ser  
 35 40 45

Val Phe Tyr His Phe Ser Val Thr Ser Phe Asp Ser Val Asp Gly Thr  
 50 55 60

Arg His Tyr Arg Val Trp Thr Ala Val Pro Asn Thr Thr Ala Pro Ala  
 65 70 75 80

Ser Gly Tyr Pro Ile Leu Tyr Met Leu Asp Gly Asn Ala Val Met Asp  
 85 90 95

Arg Leu Asp Asp Glu Leu Leu Lys Gln Leu Ser Glu Lys Thr Pro Pro  
 100 105 110

Val Ile Val Ala Val Gly Tyr Gln Thr Asn Leu Pro Phe Asp Leu Asn  
 115 120 125

Ser Arg Ala Tyr Asp Tyr Thr Pro Ala Ala Glu Ser Arg Lys Thr Asp  
 130 135 140

Leu His Ser Gly Arg Phe Ser Arg Lys Ser Gly Gly Ser Asn Asn Phe  
 145 150 155 160

Arg Gln Leu Leu Glu Thr Arg Ile Ala Pro Lys Val Glu Gln Gly Leu  
 165 170 175

Asn Ile Asp Arg Gln Arg Arg Gly Leu Trp Gly His Ser Tyr Gly Gly  
 180 185 190

Leu Phe Val Leu Asp Ser Trp Leu Ser Ser Ser Tyr Phe Arg Ser Tyr  
 195 200 205

Tyr Ser Ala Ser Pro Ser Leu Gly Arg Gly Tyr Asp Ala Leu Leu Ser  
 210 215 220



Arg Val Thr Ala Val Glu Pro Leu Gln Phe Cys Ala Lys His Leu Ala  
225 230 235 240

Ile Met Glu Gly Ser Ala Thr Gln Gly Asp Asn Arg Glu Thr His Ala  
245 250 255

Val Gly Val Leu Ser Lys Ile His Thr Thr Leu Thr Ile Leu Lys Asp  
260 265 270

Lys Gly Val Asn Ala Val Phe Trp Asp Phe Pro Asn Leu Gly His Gly  
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Pro Met Phe Asn Ala Ser Phe Arg Gln Ala Leu Leu Asp Ile Ser Gly  
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<210> 33

<211> 3292

<212> DNA

<213> *Escherichia coli*

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 Gly Arg Leu Arg Asp Ser Arg Leu Pro Met Glu Leu Trp Ala Pro Leu  
 1835 1840 1845 1850

aag gag aaa aca gat aga tat ggt ggt tta tat atc tat cga gat gga 6011  
 Lys Glu Lys Thr Asp Arg Tyr Gly Gly Leu Tyr Ile Tyr Arg Asp Gly  
 1855 1860 1865

tta aga att ttg ccc tat gga gat tca gat acg gat ttt cta aaa ata 6059  
 Leu Arg Ile Leu Pro Tyr Gly Asp Ser Asp Thr Asp Phe Leu Lys Ile  
 1870 1875 1880

gaa aag aga aga acg tta tcc gct tct gaa tat ttt ttc tca tat cga 6107  
 Glu Lys Arg Arg Thr Leu Ser Ala Ser Glu Tyr Phe Phe Ser Tyr Arg  
 1885 1890 1895

cgt ttg ttt gga gca ata gaa tta aca aaa gaa aac aat gct tca tta 6155  
 Arg Leu Phe Gly Ala Ile Glu Leu Thr Lys Glu Asn Asn Ala Ser Leu  
 1900 1905 1910

gtt gaa aaa gct ggg cga gaa gga ttc att gaa aat aag cca tat aaa 6203  
 Val Glu Lys Ala Gly Arg Glu Gly Phe Ile Glu Asn Lys Pro Tyr Lys  
 1915 1920 1925 1930

cag ttt aaa gaa atg ctt gaa aat ttc ttc atc gaa atc gca aga gat 6251  
 Gln Phe Lys Glu Met Leu Glu Asn Phe Phe Ile Glu Ile Ala Arg Asp  
 1935 1940 1945

ttc ttt aag gac gat ggc gat atg tct gaa tta ttt gtt gag aca aag 6299  
 Phe Phe Lys Asp Asp Gly Asp Met Ser Glu Leu Phe Val Glu Thr Lys  
 1950 1955 1960

caa cgt aga aat gaa gaa cat gat ttg tta tct aaa aga tct aaa caa 6347  
 Gln Arg Arg Asn Glu Glu His Asp Leu Leu Ser Lys Arg Ser Lys Gln  
 1965 1970 1975

act aaa gct aaa aaa gat aga tta aag aaa gat ctg tat gat ttt ttt 6395  
 Thr Lys Ala Lys Lys Asp Arg Leu Lys Lys Asp Leu Tyr Asp Phe Phe  
 1980 1985 1990

gat aag tta gat aat gat tac tgg aat att gaa ata aat aag cta atc 6443  
 Asp Lys Leu Asp Asn Asp Tyr Trp Asn Ile Glu Ile Asn Lys Leu Ile  
 1995 2000 2005 2010

aat aaa aac gag gaa tat ttc tcc agt aca gaa ata aca gac acc aat 6491  
 Asn Lys Asn Glu Glu Tyr Phe Ser Ser Thr Glu Ile Thr Asp Thr Asn  
 2015 2020 2025

ata gat tat gta tac aat aaa att aaa gaa caa aat gat gct atc att 6539  
 Ile Asp Tyr Val Tyr Asn Lys Ile Lys Glu Gln Asn Asp Ala Ile Ile  
 2030 2035 2040

aaa aat cta cgt aat tct gtg gat ata aag aaa ccc tct gga gtt gga 6587  
 Lys Asn Leu Arg Asn Ser Val Asp Ile Lys Lys Pro Ser Gly Val Gly  
 2045 2050 2055

tta aca aaa gag tta tct aat tta tgg gat aga tat caa ata gaa aga 6635  
 Leu Thr Lys Glu Leu Ser Asn Leu Trp Asp Arg Tyr Gln Ile Glu Arg  
 2060 2065 2070

caa aaa ata ctg tta tca cta aat gag cta aaa gat aac gtt gat aga 6683  
 Gln Lys Ile Leu Leu Ser Leu Asn Glu Leu Lys Asp Asn Val Asp Arg  
 2075 2080 2085 2090

aag ctt ata gaa ctg gat aat aaa aat aat gat ttt ctc aac tta cgg 6731  
 Lys Leu Ile Glu Leu Asp Asn Lys Asn Asn Asp Phe Leu Asn Leu Arg  
 2095 2100 2105

aag aga ctt gaa gat tct ttg aat cta caa caa agt tac tat gaa aaa 6779  
 Lys Arg Leu Glu Asp Ser Leu Asn Leu Gln Gln Ser Tyr Tyr Glu Lys  
 2110 2115 2120

gaa cta aca aag tta tat aat gac gct aaa aat gct ttg aaa gat gtg 6827  
 Glu Leu Thr Lys Leu Tyr Asn Asp Ala Lys Asn Ala Leu Lys Asp Val  
 2125 2130 2135

caa tct aaa gca aat agg tta att tct gat aat aag aaa aaa cat aag 6875  
 Gln Ser Lys Ala Asn Arg Leu Ile Ser Asp Asn Lys Lys Lys His Lys  
 2140 2145 2150

agt gaa cta aaa aac att tct tat gaa ttc caa tca act aat ctc aat 6923  
 Ser Glu Leu Lys Asn Ile Ser Tyr Glu Phe Gln Ser Thr Asn Leu Asn  
 2155 2160 2165 2170

ggc aaa gat act gcg tat ata ttg gat gta aaa aga aat cta gaa agt 6971  
 Gly Lys Asp Thr Ala Tyr Ile Leu Asp Val Lys Arg Asn Leu Glu Ser  
 2175 2180 2185

aaa att gag aat act tca aac gaa gtg att aat gaa ata aga aaa cta 7019  
 Lys Ile Glu Asn Thr Ser Asn Glu Val Ile Asn Glu Ile Arg Lys Leu  
 2190 2195 2200

acc gac cag att gca ata att agt gat agt acc act tct gaa aat tta	7067
Thr Asp Gln Ile Ala Ile Ile Ser Asp Ser Thr Thr Ser Glu Asn Leu	
2205 2210 2215	
tca tcg gct caa gta act gaa gca atc gaa act gaa ctt gaa cat tta	7115
Ser Ser Ala Gln Val Thr Glu Ala Ile Glu Thr Glu Leu Glu His Leu	
2220 2225 2230	
cga gac caa caa gca aat aac gca gag tta ata cta ctt ggc atg gct	7163
Arg Asp Gln Gln Ala Asn Asn Ala Glu Leu Ile Leu Leu Gly Met Ala	
2235 2240 2245 2250	
ctt tct gta gta cat cat gaa ttt aat ggt aat att agg gca att aga	7211
Leu Ser Val Val His His Glu Phe Asn Gly Asn Ile Arg Ala Ile Arg	
2255 2260 2265	
agt gcg cta agg gaa tta aaa gca tgg gct gac aga aat cct aag ctt	7259
Ser Ala Leu Arg Glu Leu Lys Ala Trp Ala Asp Arg Asn Pro Lys Leu	
2270 2275 2280	
gat att ata tac caa aaa atc aga act agt ttt gat cac tta gat ggt	7307
Asp Ile Ile Tyr Gln Lys Ile Arg Thr Ser Phe Asp His Leu Asp Gly	
2285 2290 2295	
tat tta aaa acc ttt aca cca ttg aca aga cgt tta agt cgc tct aaa	7355
Tyr Leu Lys Thr Phe Thr Pro Leu Thr Arg Arg Leu Ser Arg Ser Lys	
2300 2305 2310	
acc aat ata act gga act gcc att tta gaa ttt atc aga gat gta ttc	7403
Thr Asn Ile Thr Gly Thr Ala Ile Leu Glu Phe Ile Arg Asp Val Phe	
2315 2320 2325 2330	
gat gat cgt ctt gag aaa gaa gga att gaa tta ttc act acc tca aag	7451
Asp Asp Arg Leu Glu Lys Glu Gly Ile Glu Leu Phe Thr Thr Ser Lys	
2335 2340 2345	
ttt gtt aat caa gaa att gta act tac aca tca acc att tac cct gtc	7499
Phe Val Asn Gln Glu Ile Val Thr Tyr Thr Ser Thr Ile Tyr Pro Val	
2350 2355 2360	
ttt ata aat cta att gat aac gca ata tac tgg ctt ggg aaa aca act	7547
Phe Ile Asn Leu Ile Asp Asn Ala Ile Tyr Trp Leu Gly Lys Thr Thr	
2365 2370 2375	
gga gaa aaa aga ctt ata ctt gat gct act gaa aca gga ttt gtt att	7595
Gly Glu Lys Arg Leu Ile Leu Asp Ala Thr Glu Thr Gly Phe Val Ile	
2380 2385 2390	



ggt gat act ggt ccc ggt gtt tca act aga gat cga gat ata ata ttt	7643
Gly Asp Thr Gly Pro Gly Val Ser Thr Arg Asp Arg Asp Ile Ile Phe	
2395                      2400                      2405                      2410	
gat atg gga ttt aca cga aaa aca gga ggg cgt gga atg gga tta ttc	7691
Asp Met Gly Phe Thr Arg Lys Thr Gly Gly Arg Gly Met Gly Leu Phe	
2415                      2420                      2425	
att tcc aaa gag tgt tta tct cga gat gga ttt act ata aga ttg gat	7739
Ile Ser Lys Glu Cys Leu Ser Arg Asp Gly Phe Thr Ile Arg Leu Asp	
2430                      2435                      2440	
gat tac act cct gaa cag ggt gct ttc ttt att att gag cca tca gaa	7787
Asp Tyr Thr Pro Glu Gln Gly Ala Phe Phe Ile Ile Glu Pro Ser Glu	
2445                      2450                      2455	
gaa aca agt gaa tag cgatataaa taa atg aca agc tct act gat ttt	7836
Glu Thr Ser Glu                      Met Thr Ser Ser Thr Asp Phe	
2460                      2465                      2470	
cat aaa ctt tct gaa gac tgc gtt cgc cgt ttt tta cat tct gta gtt	7884
His Lys Leu Ser Glu Asp Cys Val Arg Arg Phe Leu His Ser Val Val	
2475                      2480                      2485	
gct gta gat gac aat atg tct ttt gga gct ggt agt gat act ttc cct	7932
Ala Val Asp Asp Asn Met Ser Phe Gly Ala Gly Ser Asp Thr Phe Pro	
2490                      2495                      2500	
aca gac gaa gat att aat gct tta gtt gat ccc gac gat gat cct aca	7980
Thr Asp Glu Asp Ile Asn Ala Leu Val Asp Pro Asp Asp Asp Pro Thr	
2505                      2510                      2515	
cca ata ata aca gca tca gca tcc cca agg ata gaa tca act aaa tca	8028
Pro Ile Ile Thr Ala Ser Ala Ser Pro Arg Ile Glu Ser Thr Lys Ser	
2520                      2525                      2530	
aaa gca aag gta aaa aac cat cct ttt gat tac caa gct cta gca gaa	8076
Lys Ala Lys Val Lys Asn His Pro Phe Asp Tyr Gln Ala Leu Ala Glu	
2535                      2540                      2545                      2550	
gct ttc gcc aaa gat ggt att gct tgt tgc gga tta tta gct aag agt	8124
Ala Phe Ala Lys Asp Gly Ile Ala Cys Cys Tyr Leu Leu Ala Lys Ser	
2555                      2560                      2565	
ttt aat gtt gaa gaa aga gat ata att aca gca tca tcc cac aag gca	8172
Phe Asn Val Glu Glu Arg Asp Ile Ile Thr Ala Ser Ser His Lys Ala	
2570                      2575                      2580	

gat ata aca ata ctt gac tgg gat atg caa agc gat agt ggg caa ttt	8220
Asp Ile Thr Ile Leu Asp Trp Asp Met Gln Ser Asp Ser Gly Gln Phe	
2585 2590 2595	
gct att gaa ata ata aaa tcg ata atc gtt tca gat ata aat tct gga	8268
Ala Ile Glu Ile Ile Lys Ser Ile Ile Val Ser Asp Ile Asn Ser Gly	
2600 2605 2610	
gga cgt tta cgt ctt ctt tct att tat act ggt gaa cat gtt act gct	8316
Gly Arg Leu Arg Leu Leu Ser Ile Tyr Thr Gly Glu His Val Thr Ala	
2615 2620 2625 2630	
gtt ata act aag ttg aac aat gag tta aag aaa aca tac cgt agc gta	8364
Val Ile Thr Lys Leu Asn Asn Glu Leu Lys Lys Thr Tyr Arg Ser Val	
2635 2640 2645	
ata aaa aat gat gat agt att ttt att gaa gat aac tat gca ctc gaa	8412
Ile Lys Asn Asp Asp Ser Ile Phe Ile Glu Asp Asn Tyr Ala Leu Glu	
2650 2655 2660	
caa tgg tgt ata gtt gtt att agt aaa gac gtt tat gaa aaa gat ctt	8460
Gln Trp Cys Ile Val Val Ile Ser Lys Asp Val Tyr Glu Lys Asp Leu	
2665 2670 2675	
cca aat gtg tta ata aaa aaa ttc act aac ctt aca gct ggg ttg cta	8508
Pro Asn Val Leu Ile Lys Lys Phe Thr Asn Leu Thr Ala Gly Leu Leu	
2680 2685 2690	
tcc aac gcc gca ctc tct tgc att tct gaa ata aga gaa aaa acc cat	8556
Ser Asn Ala Ala Leu Ser Cys Ile Ser Glu Ile Arg Glu Lys Thr His	
2695 2700 2705 2710	
ggg ata tta aca aaa tat aat aat aaa tta gac act gca tat gtt tcc	8604
Gly Ile Leu Thr Lys Tyr Asn Asn Lys Leu Asp Thr Ala Tyr Val Ser	
2715 2720 2725	
cac atc tta aat tta ata aaa tcc aag gag tca agg gca tat gct tat	8652
His Ile Leu Asn Leu Ile Lys Ser Lys Glu Ser Arg Ala Tyr Ala Tyr	
2730 2735 2740	
gaa aat gct cat gat tat gca gta gat tta att tct gaa gaa ata aga	8700
Glu Asn Ala His Asp Tyr Ala Val Asp Leu Ile Ser Glu Glu Ile Arg	
2745 2750 2755	
tca ata ttg caa ata agt gaa aac tta aag aaa tct cta agc aaa aac	8748
Ser Ile Leu Gln Ile Ser Glu Asn Leu Lys Lys Ser Leu Ser Lys Asn	
2760 2765 2770	

tcc tta tcc cat tgg cct att ttt cac tat gca aaa aat ggt tgt aag 8796  
 Ser Leu Ser His Trp Pro Ile Phe His Tyr Ala Lys Asn Gly Cys Lys  
 2775 2780 2785 2790

aat ttt cta tta act gga aaa aaa caa aaa gac tta tca gta gaa cat 8844  
 Asn Phe Leu Leu Thr Gly Lys Lys Gln Lys Asp Leu Ser Val Glu His  
 2795 2800 2805

cta agg aat ata ctc tct gct gat tct tta gaa gaa att caa cac gct 8892  
 Leu Arg Asn Ile Leu Ser Ala Asp Ser Leu Glu Glu Ile Gln His Ala  
 2810 2815 2820

att gaa cac gca tct tta ggt aaa aag gaa tac tta agc caa gat ggt 8940  
 Ile Glu His Ala Ser Leu Gly Lys Lys Glu Tyr Leu Ser Gln Asp Gly  
 2825 2830 2835

gaa gaa gat aaa aag tta atg caa tta tgc tct ctg gaa atc acg cgc 8988  
 Glu Glu Asp Lys Lys Leu Met Gln Leu Cys Ser Leu Glu Ile Thr Arg  
 2840 2845 2850

agg agt tta aga tat cat tct cat ata gat aat gtg tcc tta aaa caa 9036  
 Arg Ser Leu Arg Tyr His Ser His Ile Asp Asn Val Ser Leu Lys Gln  
 2855 2860 2865 2870

gga act tta ctt tta gat gca tat aat ttt gtc tat cta tgc ata caa 9084  
 Gly Thr Leu Leu Leu Asp Ala Tyr Asn Phe Val Tyr Leu Cys Ile Gln  
 2875 2880 2885

cca tta tgt gat agc gtc aga ttg cat gaa aaa gcc gat ttt tta ttc 9132  
 Pro Leu Cys Asp Ser Val Arg Leu His Glu Lys Ala Asp Phe Leu Phe  
 2890 2895 2900

ctc agg gga aca ctg gac gat aat aat tac aat ttg tta atc gaa gat 9180  
 Leu Arg Gly Thr Leu Asp Asp Asn Asn Tyr Asn Leu Leu Ile Glu Asp  
 2905 2910 2915

gaa tat ggc ggt ttt tat aaa att aaa atg ccg gca aaa gct tct aat 9228  
 Glu Tyr Gly Gly Phe Tyr Lys Ile Lys Met Pro Ala Lys Ala Ser Asn  
 2920 2925 2930

att att tca ttt tca ttt gga gtc gaa aat gga aac ggt gtc atc ata 9276  
 Ile Ile Ser Phe Ser Phe Gly Val Glu Asn Gly Asn Gly Val Ile Ile  
 2935 2940 2945 2950

ggg aaa aag aac aat cta gtt aat act gac tat atc tca ttc gtt cct 9324  
 Gly Lys Lys Asn Asn Leu Val Asn Thr Asp Tyr Ile Ser Phe Val Pro  
 2955 2960 2965

tta ctc gtt gaa aaa ata tct act cca aaa gta ttg aaa tgg atc ggg 9372  
 Leu Leu Val Glu Lys Ile Ser Thr Pro Lys Val Leu Lys Trp Ile Gly  
 2970 2975 2980

gaa ata aaa aca acg tac gcg caa aaa ata aca act gat att gtt gct 9420  
 Glu Ile Lys Thr Thr Tyr Ala Gln Lys Ile Thr Thr Asp Ile Val Ala  
 2985 2990 2995

aat ctg tca aga ata ggt tta gat caa cat gag tgg tta cga ata aaa 9468  
 Asn Leu Ser Arg Ile Gly Leu Asp Gln His Glu Trp Leu Arg Ile Lys  
 3000 3005 3010

tca aaa gat ata taaatgatta tatatgccgt cggtttataa aaactggcgg 9520  
 Ser Lys Asp Ile  
 3015

catgtatatc tagttagtcc atcatagaag tcaagaaatt tagtttgccc tatatcttat 9580

agaaaatata ttttatatgc ttaaaaaaca ccattctttct aagatggcat ttatgtgctt 9640

tgtttcgac aattacaact gatattattac catattgatt aattttatgt tattttaccaa 9700

agtaacggca tcttaataata tcgtcataat atagtgcgcg ttctgactct aatactgaaa 9760

aattttattg ttctatttta cacttactgc aaatagcatc cagtttatca tatagtgtcg 9820

catcaattgg cgacg atg tca tca cgc caa atc ctt gag cat tat aat gct 9871  
 Met Ser Ser Arg Gln Ile Leu Glu His Tyr Asn Ala  
 3020 3025 3030

cta aca tat ccc cta cat caa tca atc ttg ttg cag ata atg act tcg 9919  
 Leu Thr Tyr Pro Leu His Gln Ser Ile Leu Leu Gln Ile Met Thr Ser  
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aat ttg tta tca gtt tgc act gga aaa tcc att tac gag gat atc tcc 9967  
 Asn Leu Leu Ser Val Cys Thr Gly Lys Ser Ile Tyr Glu Asp Ile Ser  
 3050 3055 3060

ggc agt tct tgg aat atc ata cac ttc aat atc cct ctc ccc atc tct 10015  
 Gly Ser Ser Trp Asn Ile Ile His Phe Asn Ile Pro Leu Pro Ile Ser  
 3065 3070 3075

aga gcg aga ctt tcc ata ttt tct tat tgt gtc aga att aaa cct tgg 10063  
 Arg Ala Arg Leu Ser Ile Phe Ser Tyr Cys Val Arg Ile Lys Pro Trp  
 3080 3085 3090

atg agt atg gat tac atg taaccggctc atttaaaccg tctggtctgt 10111  
 Met Ser Met Asp Tyr Met

3095

3100

ttcctccggt ttacaaaaa ta atg tcc atc att ttt aat gga cac tat cgt 10163  
 Met Ser Ile Ile Phe Asn Gly His Tyr Arg  
 3105 3110

atg aaa cac cgg act tgg atc act gaa gct tta cgt ctt cac ttt gaa 10211  
 Met Lys His Arg Thr Trp Ile Thr Glu Ala Leu Arg Leu His Phe Glu  
 3115 3120 3125

gaa cat tta ccc cag gtt gtg gtc ggg cgt cgc ctg ggc gta cca aaa 10259  
 Glu His Leu Pro Gln Val Val Val Gly Arg Arg Leu Gly Val Pro Lys  
 3130 3135 3140

tca aca gct tgt ggt atg ttc gtg cgc ttt cgc aaa gct ggc ttt tca 10307  
 Ser Thr Ala Cys Gly Met Phe Val Arg Phe Arg Lys Ala Gly Phe Ser  
 3145 3150 3155

tgg cct ctg ccc gca ggt atg tcg gag cgg gag ctt gat ggc cgt ctt 10355  
 Trp Pro Leu Pro Ala Gly Met Ser Glu Arg Glu Leu Asp Gly Arg Leu  
 3160 3165 3170

tac ggg agt acc tcc aca gta cct gtc gta ctt tgt agt gga tcg gta 10403  
 Tyr Gly Ser Thr Ser Thr Val Pro Val Val Leu Cys Ser Gly Ser Val  
 3175 3180 3185 3190

att cag gac acc tcg aaa tcc tgt taatgttaaa acagtgttaaa tgagggtgatg 10457  
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 3195

c atg atc aaa act cgt cgg act aaa cgt acc ttt tcc ccg gag ttc aag 10506  
 Met Ile Lys Thr Arg Thr Lys Arg Thr Phe Ser Pro Glu Phe Lys  
 3200 3205 3210

ctt gaa gct ttc gag cag gtg gtg gtt aaa tac cag cgt gat gtc aga 10554  
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 3215 3220 3225 3230

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 3235 3240 3245

ata cgg ttg tat aag cag gaa ctt cag ggt att gag cca gct ggt aat 10650  
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 3250 3255 3260

gct att acc cct gaa caa cgc gaa att cag cag ctt aaa gcg cag ata 10698  
 Ala Ile Thr Pro Glu Gln Arg Glu Ile Gln Gln Leu Lys Ala Gln Ile

3265

3270

3275

aag cgc gtt gag atg gaa aaa gaa ata cta aag cag gct gcc gtg ctg 10746  
 Lys Arg Val Glu Met Glu Lys Glu Ile Leu Lys Gln Ala Val Leu  
 3280 3285 3290

atg agc gaa atc ccc ggg aag ctg tcg cgc taatcacaca gctgaaagca 10796  
 Met Ser Glu Ile Pro Gly Lys Leu Ser Arg  
 3295 3300

aagtggccag tgtgggttat ttgtcattta ttcggtatta accgtagcgt ttattacgcg 10856

cagggtgaagc gtcctgttaa tgtgcaaaga attgaattac gaagccgggt gagggctttc 10916

catgctctca gtcgtggcgc agccgggtag ccgggcaatc agtcagatgt tgcgccagag 10976

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tcgacagccg gttaaacatc acaaccgggt aaacgaagac aaaagtccgc cattgccaaa 11096

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<211> 366

<212> PRT

<213> Escherichia coli

<400> 35

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 20 25 30

Glu Leu Ala Asn Ile Ile Gly His His Ala Gly Ile Asp Asp Asn Thr  
 35 40 45

Ala Ala Lys Ala Ile Ala His Ala Ile Leu Gly Gly Val Thr Ala Ala  
 50 55 60

Leu Gln Gly Asn Ser Ala Ala Gly Ala Ile Gly Ala Gly Thr Gly  
 65 70 75 80

Glu Val Ile Ala Ser Ala Ile Ala Lys Ser Leu Tyr Pro Gly Val Asp  
 85 90 95

Pro Ser Lys Leu Thr Glu Asp Gln Lys Gln Thr Val Ser Thr Leu Ala  
 100 105 110

Thr Leu Ser Ala Gly Met Ala Gly Gly Ile Ala Ser Gly Asp Val Ala  
 115 120 125

Gly Ala Ala Ala Gly Ala Gly Ala Gly Lys Asn Val Val Glu Asn Asn  
 130 135 140

Ala Leu Ser Leu Val Ala Arg Gly Cys Ala Val Ala Ala Pro Cys Arg  
 145 150 155 160

Thr Lys Val Ala Glu Gln Leu Leu Glu Ile Gly Ala Lys Ala Gly Met  
 165 170 175

Ala Gly Leu Ala Gly Ala Ala Val Lys Asp Met Ala Asp Arg Met Thr  
 180 185 190

Ser Asp Glu Leu Glu His Leu Ile Thr Leu Gln Met Met Gly Asn Asp  
 195 200 205

Glu Ile Thr Thr Lys Tyr Leu Ser Ser Leu His Asp Lys Tyr Gly Ser  
 210 215 220

Gly Ala Ala Ser Asn Pro Asn Ile Gly Lys Asp Leu Thr Asp Ala Glu  
 225 230 235 240

Lys Val Glu Leu Gly Gly Ser Gly Ser Gly Thr Gly Thr Pro Pro Pro  
 245 250 255

Ser Glu Asn Asp Pro Lys Gln Gln Asn Glu Lys Thr Val Asp Lys Leu  
 260 265 270

Asn Gln Lys Gln Glu Ser Ala Ile Lys Lys Ile Asp Asn Thr Ile Lys  
 275 280 285

Asn Ala Leu Lys Asp His Asp Ile Ile Gly Thr Leu Lys Asp Met Asp  
 290 295 300

Gly Lys Pro Val Pro Lys Glu Asn Gly Gly Tyr Trp Asp His Met Gln  
 305 310 315 320

Glu Met Gln Asn Thr Leu Arg Gly Leu Arg Asn His Ala Asp Thr Leu  
 325 330 335

Lys Asn Val Asn Asn Pro Glu Ala Gln Ala Ala Tyr Gly Arg Ala Thr  
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Asp Ala Ile Asn Lys Ile Glu Ser Ala Leu Lys Gly Tyr Gly  
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<213> Escherichia coli

<400> 36

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Val Pro Leu Glu Lys Leu Thr Val Glu Asp Leu Cys Arg Ala Ile Arg  
 35 40 45

Gln Asn Leu Cys Ile Asp Gln Leu Met Pro Arg Val Leu Glu Val Leu  
 50 55 60

Thr Lys Glu Pro Leu Ala Gly Glu Tyr Tyr Asp Gly Glu Leu Ile Ala  
 65 70 75 80

Ala Leu Ser Thr Ile Lys Gly Glu Asp Leu Lys Asp Gln Lys Ser Thr  
 85 90 95

Phe Thr Gln Ile Arg Gln Leu Ile Asn Gln Leu Glu Pro Ser Asp Ile  
 100 105 110

Asn Asp Asp Leu Arg Lys Asp Ile Leu Lys Ile Asn Gln Ile Ile Val  
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<212> PRT

<213> Escherichia coli

<400> 37

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 20 25 30



Glu Asn Ala Leu Arg Ala Val Ser Leu Gly Arg Lys Asn Phe Leu Phe  
 35 40 45

Phe Gly Ser Asp His Gly Gly Glu Arg Gly Ala Leu Leu Tyr Ser Leu  
 50 55 60

Ile Gly Thr Cys Lys Leu Asn Asp Val Asp Pro Glu Ser Tyr Leu Arg  
 65 70 75 80

His Val Leu Ala Val Ile Ala Asp Trp Pro Val Asn Arg Val Ser Glu  
 85 90 95

Leu Leu Pro Trp Arg Ile Ala Leu Pro Ala Glu  
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<400> 38

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 1 5 10 15

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 20 25 30

Ile Asn Leu Asn Pro Val Ser Ala Leu Ser Ala Leu Asp Asn Pro Phe  
 35 40 45

Arg Ser Ala Asp Asn Ala Thr Gly Arg Ile Thr Ser Ser Ile Gln Pro  
 50 55 60

Ala Val Gln Cys Ala Ala Ala Ala Thr Glu Gly Ser Cys Pro Arg  
 65 70 75 80

Gln Ser Pro Cys Ser Gly  
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<211> 111

<212> PRT

<213> Escherichia coli

<400> 39

Met Val Asp Asn Trp Gln Lys Ser Val Arg Ser Arg Ala Leu Pro Glu

1                      5                      10                      15  
 Glu Ala Met Thr Gly Trp Asn Glu Gly Met Ile Arg Leu Gln Gln Leu  
                     20                      25                      30  
 Ala Glu Arg Leu Asn Arg Gln Asp Glu Gln Arg Gly Lys Tyr Met Thr  
                     35                      40                      45  
 Val Ser Glu Leu Lys Thr Glu Val Phe Gly Ile Met Gln Ala Phe Asn  
                     50                      55                      60  
 Arg His Ile Pro Ala Glu Glu Gln Leu Arg Arg Tyr Gly Glu Val Arg  
                     65                      70                      75                      80  
 Asn Gln Asn Gly Ser Glu Gln Gln Gln Lys Gln Ala Glu Met Ala Leu  
                     85                      90                      95  
 Asn Gln Leu Ile Asn Arg Tyr Gln Met Ile Arg Ala Gly Lys Gln  
                     100                      105                      110  
  
 <210> 40  
 <211> 143  
 <212> PRT  
 <213> *Escherichia coli*  
  
 <400> 40  
 Met Val Gly Cys Ala Trp Leu Ala Glu Gln Ala Phe Ser Asp His Ala  
                     1                      5                      10                      15  
 Leu Ser Pro His Ser Ala Trp Pro Tyr Ser Ala Ser Arg Asp Ala Gly  
                     20                      25                      30  
 Leu Ala Asp Thr Gly Ala Gly Gly Tyr Pro Thr Cys Lys Gln Arg Trp  
                     35                      40                      45  
 Ala Asp Asp Thr Val Gly Leu Lys Ala Arg Leu Leu Gln Leu Pro Ala  
                     50                      55                      60  
 Leu Asp Ile Trp Thr Ala Phe Lys Lys Ile Asp Gln Ser Gln Val Val  
                     65                      70                      75                      80  
 Tyr Glu Glu Ala Val Leu Arg Ser Arg Val Ser Glu Arg Asn Met Gln  
                     85                      90                      95  
 Val Ser Gln Asn Gly Arg Val Tyr Pro Ser Tyr Gly Gly Asn Val Asp  
                     100                      105                      110

Gly Thr Val Ala Asn Ala Ala Thr Arg Leu Ala Ser Gly Ala Arg Asn  
 115 120 125

Ile Leu Gly Ser Ile Ala Ala Cys Thr Ala Phe Asp Ser Val Arg  
 130 135 140

<210> 41

<211> 118

<212> PRT

<213> Escherichia coli

<400> 41

Met Val Gln Ala Gln Leu Gln Ile Ala Leu Val Ile Cys Ile Pro Leu  
 1 5 10 15

Ile Thr Leu Cys Ser Ala Trp Asp Val Lys Val Val Met Thr Leu Thr  
 20 25 30

Phe Val Gln Phe Ala Leu Phe Phe Leu Thr Phe Trp Trp Glu Leu Ala  
 35 40 45

Arg Trp Leu Asp Ser Trp Leu Leu Asp Val Leu Tyr Asn Ser Asp Thr  
 50 55 60

His Ser Ser Trp Asn Leu Ala Gly Ile Gln Asn Thr Gln Asp Asp Val  
 65 70 75 80

Ile Ile Asn Leu Val Met Arg Leu Met Phe Leu Val Leu Pro Thr Phe  
 85 90 95

Trp Leu Gly Ala Met Thr Trp Ala Gly Val Arg Val Gly Val Ala Leu  
 100 105 110

Asn Gly Ala Leu Ala Gly  
 115

<210> 42

<211> 81

<212> PRT

<213> Escherichia coli

<400> 42

Met Lys Tyr Leu Phe Phe Glu Asn Ile His Ser Ile Phe Leu Thr Phe  
 1 5 10 15

Ser Leu Phe Arg Thr Ser Val Ser Pro Asp Phe Pro Met Ile Phe Ala

[illegible]

Ala

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<210> 43
<211> 348
<212> PRT
<213> Escherichia coli
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<400> 43

Leu	Ile	Val	Ile	Asp	Phe	Phe	Cys	Gly	Cys	Gly	Gly	Ala	Ser	Glu	Gly
1				5					10					15	
Leu	Arg	Gln	Ala	Gly	Phe	Asp	Ile	Glu	Leu	Gly	Leu	Asp	Ile	Asp	Gln
			20					25					30		
Gln	Ala	Ser	Glu	Thr	Phe	Lys	Ala	Asn	Phe	Pro	Asp	Ala	Lys	Phe	Ile
		35					40						45		
Gln	Asp	Asp	Ile	Arg	Lys	Ile	Glu	Pro	Gln	Asp	Ile	Ser	Asp	Ile	Ile
	50					55					60				
Asp	Ile	Lys	Ala	Lys	Arg	Pro	Leu	Leu	Leu	Ser	Ala	Cys	Ala	Pro	Cys
65					70					75					80
Gln	Pro	Phe	Ser	Gln	Gln	Asn	Lys	Asn	Lys	Thr	Ser	Asp	Asp	Ser	Arg
				85					90					95	
Arg	Asn	Leu	Leu	Asn	Glu	Thr	His	Arg	Phe	Ile	Arg	Glu	Leu	Leu	Pro
		100						105					110		
Glu	Tyr	Ile	Met	Leu	Glu	Asn	Val	Pro	Gly	Met	Gln	Lys	Ile	Asp	Glu
	115						120					125			
Glu	Lys	Glu	Gly	Pro	Phe	Gln	Glu	Phe	Ile	Lys	Leu	Leu	Lys	Glu	Leu
	130					135					140				
Glu	Tyr	Asn	Tyr	Ile	Ser	Phe	Ile	Ala	Asn	Ala	Glu	Asn	Tyr	Gly	Ile

145                      150                      155                      160  
 Pro Gln Arg Arg Lys Arg Leu Val Leu Leu Ala Ser Arg Val Gly Lys  
                          165                      170                      175  
 Val Thr Leu Pro Glu Ile Thr His Gly Lys Asn Lys Ile Pro Phe Lys  
                          180                      185                      190  
 Thr Val Arg Asp Tyr Ile Gln Asp Phe Thr Lys Leu Cys Ser Gly Glu  
                          195                      200                      205  
 Thr Asp Pro Lys Asp Pro Leu His Arg Ala Gly Thr Leu Ser Pro Leu  
                          210                      215                      220  
 Asn Leu Lys Arg Ile Met His Thr Pro Glu Gly Gly Asp Arg Arg Asn  
                          225                      230                      235                      240  
 Trp Pro Glu Glu Leu Val Asn Lys Cys His Lys Asn Tyr Asp Gly His  
                          245                      250                      255  
 Thr Asp Thr Tyr Gly Arg Met Ser Trp Asp Lys Pro Ala Pro Thr Leu  
                          260                      265                      270  
 Thr Thr Lys Cys Asn Ser Tyr Ser Asn Gly Arg Phe Gly His Pro Asp  
                          275                      280                      285  
 Pro Thr Gln His Arg Ala Ile Ser Ile Arg Glu Ala Ser Arg Leu Gln  
                          290                      295                      300  
 Thr Phe Pro Leu Ser Tyr Val Phe Lys Gly Ser Leu Asn Ser Met Ala  
                          305                      310                      315                      320  
 Lys Gln Ile Gly Asn Ala Val Pro Cys Glu Leu Ala Arg Leu Phe Gly  
                          325                      330                      335  
 Leu His Leu Ile Glu Asn Cys Thr Asn Lys Asp Ser  
                          340                      345

&lt;210&gt; 44

&lt;211&gt; 974

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 44

Met Leu Gly Arg Gln Gln Ile Ala Gly Ile Pro Thr Ala Leu Ser Glu  
     1                      5                      10                      15

Leu Phe Lys Asn Ala His Asp Ala Tyr Ala Asp Asn Val Glu Val Asp  
 20 25 30

Phe Phe Arg Lys Glu Asn Leu Leu Ile Leu Arg Asp Asp Gly Leu Gly  
 35 40 45

Met Thr Thr Asp Glu Phe Glu Glu Arg Trp Leu Thr Ile Gly Thr Ser  
 50 55 60

Ser Lys Leu Ile Asp Asp Asp Ala Ile Asn Lys Pro Ala Val Asp Ser  
 65 70 75 80

Asn Lys Ala Phe Arg Pro Ile Met Gly Glu Lys Gly Ile Gly Arg Leu  
 85 90 95

Ser Ile Ala Ala Ile Gly Pro Gln Val Leu Val Leu Thr Arg Ala Lys  
 100 105 110

Arg Asp Asn Glu Leu Lys Pro Leu Val Ala Ala Phe Val Asn Trp Ser  
 115 120 125

Leu Phe Ala Ile Pro Ser Leu Asp Leu Asp Asp Ile Glu Ile Pro Ile  
 130 135 140

Arg Thr Ile Ile Asn Asp Glu Cys Phe Thr Lys Lys Thr Leu Asp Glu  
 145 150 155 160

Met Ile Glu Gln Ala Arg Asn Asn Leu Asp Ser Leu Ser His Lys Ile  
 165 170 175

Ser Lys Ser Lys Val Ser Gln Ile Asn Thr Gln Leu Ser Ser Phe Glu  
 180 185 190

Phe Asp Pro Ile Leu Trp Glu Lys Lys Leu Gly Gly Leu Arg Leu Ser  
 195 200 205

Gly Asp Gly His Gly Thr His Phe Ile Ile Met Pro Thr Glu Glu Ile  
 210 215 220

Leu Ile Asp Asp Ile Ser Thr Ser Asp Ser Asn Lys Thr Ser Glu Gln  
 225 230 235 240

Ser Ser Arg Leu Glu Lys Ala Leu Leu Gly Phe Thr Asn Thr Met Tyr  
 245 250 255

Ser Asp Ser Asn Pro Pro Ile Ile Ala Arg Phe Arg Asp Tyr Leu Glu  
 260 265 270

Asp Gly Glu Cys Ile Asp Arg Ile Ser Glu Ser Ile Phe Phe Thr Pro  
 275 280 285

Gln Glu Phe Asn Leu Ala Asp His His Ile Glu Gly Trp Phe Asn Glu  
 290 295 300

Phe Gly Gln Phe Ser Gly Thr Val Ser Val Tyr Gly Glu Glu Pro Ile  
 305 310 315 320

His His Val Val Thr Trp Lys Asn Asn Asn Gln Leu Thr Gln Cys Gly  
 325 330 335

Pro Phe Lys Ile Lys Leu Ala Tyr Ile His Gly Arg Leu Arg Asp Ser  
 340 345 350

Arg Leu Pro Met Glu Leu Trp Ala Pro Leu Lys Glu Lys Thr Asp Arg  
 355 360 365

Tyr Gly Gly Leu Tyr Ile Tyr Arg Asp Gly Leu Arg Ile Leu Pro Tyr  
 370 375 380

Gly Asp Ser Asp Thr Asp Phe Leu Lys Ile Glu Lys Arg Arg Thr Leu  
 385 390 395 400

Ser Ala Ser Glu Tyr Phe Phe Ser Tyr Arg Arg Leu Phe Gly Ala Ile  
 405 410 415

Glu Leu Thr Lys Glu Asn Asn Ala Ser Leu Val Glu Lys Ala Gly Arg  
 420 425 430

Glu Gly Phe Ile Glu Asn Lys Pro Tyr Lys Gln Phe Lys Glu Met Leu  
 435 440 445

Glu Asn Phe Phe Ile Glu Ile Ala Arg Asp Phe Phe Lys Asp Asp Gly  
 450 455 460

Asp Met Ser Glu Leu Phe Val Glu Thr Lys Gln Arg Arg Asn Glu Glu  
 465 470 475 480

His Asp Leu Leu Ser Lys Arg Ser Lys Gln Thr Lys Ala Lys Lys Asp  
 485 490 495

Arg Leu Lys Lys Asp Leu Tyr Asp Phe Phe Asp Lys Leu Asp Asn Asp  
 500 505 510

Tyr Trp Asn Ile Glu Ile Asn Lys Leu Ile Asn Lys Asn Glu Glu Tyr  
 515 520 525

Phe Ser Ser Thr Glu Ile Thr Asp Thr Asn Ile Asp Tyr Val Tyr Asn  
 530 535 540

Lys Ile Lys Glu Gln Asn Asp Ala Ile Ile Lys Asn Leu Arg Asn Ser  
 545 550 555 560

Val Asp Ile Lys Lys Pro Ser Gly Val Gly Leu Thr Lys Glu Leu Ser  
 565 570 575

Asn Leu Trp Asp Arg Tyr Gln Ile Glu Arg Gln Lys Ile Leu Leu Ser  
 580 585 590

Leu Asn Glu Leu Lys Asp Asn Val Asp Arg Lys Leu Ile Glu Leu Asp  
 595 600 605

Asn Lys Asn Asn Asp Phe Leu Asn Leu Arg Lys Arg Leu Glu Asp Ser  
 610 615 620

Leu Asn Leu Gln Gln Ser Tyr Tyr Glu Lys Glu Leu Thr Lys Leu Tyr  
 625 630 635 640

Asn Asp Ala Lys Asn Ala Leu Lys Asp Val Gln Ser Lys Ala Asn Arg  
 645 650 655

Leu Ile Ser Asp Asn Lys Lys Lys His Lys Ser Glu Leu Lys Asn Ile  
 660 665 670

Ser Tyr Glu Phe Gln Ser Thr Asn Leu Asn Gly Lys Asp Thr Ala Tyr  
 675 680 685

Ile Leu Asp Val Lys Arg Asn Leu Glu Ser Lys Ile Glu Asn Thr Ser  
 690 695 700

Asn Glu Val Ile Asn Glu Ile Arg Lys Leu Thr Asp Gln Ile Ala Ile  
 705 710 715 720

Ile Ser Asp Ser Thr Thr Ser Glu Asn Leu Ser Ser Ala Gln Val Thr  
 725 730 735

Glu Ala Ile Glu Thr Glu Leu Glu His Leu Arg Asp Gln Gln Ala Asn  
 740 745 750

Asn Ala Glu Leu Ile Leu Leu Gly Met Ala Leu Ser Val Val His His  
 755 760 765

Glu Phe Asn Gly Asn Ile Arg Ala Ile Arg Ser Ala Leu Arg Glu Leu  
 770 775 780



Lys Ala Trp Ala Asp Arg Asn Pro Lys Leu Asp Ile Ile Tyr Gln Lys  
785 790 795 800

Ile Arg Thr Ser Phe Asp His Leu Asp Gly Tyr Leu Lys Thr Phe Thr  
805 810 815

Pro Leu Thr Arg Arg Leu Ser Arg Ser Lys Thr Asn Ile Thr Gly Thr  
820 825 830

Ala Ile Leu Glu Phe Ile Arg Asp Val Phe Asp Asp Arg Leu Glu Lys  
835 840 845

Glu Gly Ile Glu Leu Phe Thr Thr Ser Lys Phe Val Asn Gln Glu Ile  
850 855 860

Val Thr Tyr Thr Ser Thr Ile Tyr Pro Val Phe Ile Asn Leu Ile Asp  
865 870 875 880

Asn Ala Ile Tyr Trp Leu Gly Lys Thr Thr Gly Glu Lys Arg Leu Ile  
885 890 895

Leu Asp Ala Thr Glu Thr Gly Phe Val Ile Gly Asp Thr Gly Pro Gly  
900 905 910

Val Ser Thr Arg Asp Arg Asp Ile Ile Phe Asp Met Gly Phe Thr Arg  
915 920 925

Lys Thr Gly Gly Arg Gly Met Gly Leu Phe Ile Ser Lys Glu Cys Leu  
930 935 940

Ser Arg Asp Gly Phe Thr Ile Arg Leu Asp Asp Tyr Thr Pro Glu Gln  
945 950 955 960

Gly Ala Phe Phe Ile Ile Glu Pro Ser Glu Glu Thr Ser Glu  
965 970

<210> 45

<211> 555

<212> PRT

<213> Escherichia coli

<400> 45

Met Thr Ser Ser Thr Asp Phe His Lys Leu Ser Glu Asp Cys Val Arg  
1 5 10 15

Arg Phe Leu His Ser Val Val Ala Val Asp Asp Asn Met Ser Phe Gly  
20 25 30

Ala Gly Ser Asp Thr Phe Pro Thr Asp Glu Asp Ile Asn Ala Leu Val  
 35 40 45

Asp Pro Asp Asp Asp Pro Thr Pro Ile Ile Thr Ala Ser Ala Ser Pro  
 50 55 60

Arg Ile Glu Ser Thr Lys Ser Lys Ala Lys Val Lys Asn His Pro Phe  
 65 70 75 80

Asp Tyr Gln Ala Leu Ala Glu Ala Phe Ala Lys Asp Gly Ile Ala Cys  
 85 90 95

Cys Gly Leu Leu Ala Lys Ser Phe Asn Val Glu Glu Arg Asp Ile Ile  
 100 105 110

Thr Ala Ser Ser His Lys Ala Asp Ile Thr Ile Leu Asp Trp Asp Met  
 115 120 125

Gln Ser Asp Ser Gly Gln Phe Ala Ile Glu Ile Ile Lys Ser Ile Ile  
 130 135 140

Val Ser Asp Ile Asn Ser Gly Gly Arg Leu Arg Leu Leu Ser Ile Tyr  
 145 150 155 160

Thr Gly Glu His Val Thr Ala Val Ile Thr Lys Leu Asn Asn Glu Leu  
 165 170 175

Lys Lys Thr Tyr Arg Ser Val Ile Lys Asn Asp Asp Ser Ile Phe Ile  
 180 185 190

Glu Asp Asn Tyr Ala Leu Glu Gln Trp Cys Ile Val Val Ile Ser Lys  
 195 200 205

Asp Val Tyr Glu Lys Asp Leu Pro Asn Val Leu Ile Lys Lys Phe Thr  
 210 215 220

Asn Leu Thr Ala Gly Leu Leu Ser Asn Ala Ala Leu Ser Cys Ile Ser  
 225 230 235 240

Glu Ile Arg Glu Lys Thr His Gly Ile Leu Thr Lys Tyr Asn Asn Lys  
 245 250 255

Leu Asp Thr Ala Tyr Val Ser His Ile Leu Asn Leu Ile Lys Ser Lys  
 260 265 270

Glu Ser Arg Ala Tyr Ala Tyr Glu Asn Ala His Asp Tyr Ala Val Asp  
 275 280 285

Leu Ile Ser Glu Glu Ile Arg Ser Ile Leu Gln Ile Ser Glu Asn Leu  
 290 295 300

Lys Lys Ser Leu Ser Lys Asn Ser Leu Ser His Trp Pro Ile Phe His  
 305 310 315 320

Tyr Ala Lys Asn Gly Cys Lys Asn Phe Leu Leu Thr Gly Lys Lys Gln  
 325 330 335

Lys Asp Leu Ser Val Glu His Leu Arg Asn Ile Leu Ser Ala Asp Ser  
 340 345 350

Leu Glu Glu Ile Gln His Ala Ile Glu His Ala Ser Leu Gly Lys Lys  
 355 360 365

Glu Tyr Leu Ser Gln Asp Gly Glu Glu Asp Lys Lys Leu Met Gln Leu  
 370 375 380

Cys Ser Leu Glu Ile Thr Arg Arg Ser Leu Arg Tyr His Ser His Ile  
 385 390 395 400

Asp Asn Val Ser Leu Lys Gln Gly Thr Leu Leu Leu Asp Ala Tyr Asn  
 405 410 415

Phe Val Tyr Leu Cys Ile Gln Pro Leu Cys Asp Ser Val Arg Leu His  
 420 425 430

Glu Lys Ala Asp Phe Leu Phe Leu Arg Gly Thr Leu Asp Asp Asn Asn  
 435 440 445

Tyr Asn Leu Leu Ile Glu Asp Glu Tyr Gly Gly Phe Tyr Lys Ile Lys  
 450 455 460

Met Pro Ala Lys Ala Ser Asn Ile Ile Ser Phe Ser Phe Gly Val Glu  
 465 470 475 480

Asn Gly Asn Gly Val Ile Ile Gly Lys Lys Asn Asn Leu Val Asn Thr  
 485 490 495

Asp Tyr Ile Ser Phe Val Pro Leu Leu Val Glu Lys Ile Ser Thr Pro  
 500 505 510

Lys Val Leu Lys Trp Ile Gly Glu Ile Lys Thr Thr Tyr Ala Gln Lys  
 515 520 525

Ile Thr Thr Asp Ile Val Ala Asn Leu Ser Arg Ile Gly Leu Asp Gln  
 530 535 540

His Glu Trp Leu Arg Ile Lys Ser Lys Asp Ile  
 545 550 555

<210> 46

<211> 82

<212> PRT

<213> Escherichia coli

<400> 46

Met Ser Ser Arg Gln Ile Leu Glu His Tyr Asn Ala Leu Thr Tyr Pro  
 1 5 10 15

Leu His Gln Ser Ile Leu Leu Gln Ile Met Thr Ser Asn Leu Leu Ser  
 20 25 30

Val Cys Thr Gly Lys Ser Ile Tyr Glu Asp Ile Ser Gly Ser Ser Trp  
 35 40 45

Asn Ile Ile His Phe Asn Ile Pro Leu Pro Ile Ser Arg Ala Arg Leu  
 50 55 60

Ser Ile Phe Ser Tyr Cys Val Arg Ile Lys Pro Trp Met Ser Met Asp  
 65 70 75 80

Tyr Met

<210> 47

<211> 98

<212> PRT

<213> Escherichia coli

<400> 47

Met Ser Ile Ile Phe Asn Gly His Tyr Arg Met Lys His Arg Thr Trp  
 1 5 10 15

Ile Thr Glu Ala Leu Arg Leu His Phe Glu Glu His Leu Pro Gln Val  
 20 25 30

Val Val Gly Arg Arg Leu Gly Val Pro Lys Ser Thr Ala Cys Gly Met  
 35 40 45

Phe Val Arg Phe Arg Lys Ala Gly Phe Ser Trp Pro Leu Pro Ala Gly  
 50 55 60

Met Ser Glu Arg Glu Leu Asp Gly Arg Leu Tyr Gly Ser Thr Ser Thr

65

70

75

80

Val Pro Val Val Leu Cys Ser Gly Ser Val Ile Gln Asp Thr Ser Lys  
                   85                                  90                                  95

Ser Cys

&lt;210&gt; 48

&lt;211&gt; 106

&lt;212&gt; PRT

&lt;213&gt; Escherichia coli

&lt;400&gt; 48

Met Ile Lys Thr Arg Arg Thr Lys Arg Thr Phe Ser Pro Glu Phe Lys  
   1                                  5                                  10                                  15

Leu Glu Ala Phe Glu Gln Val Val Val Lys Tyr Gln Arg Asp Val Arg  
                   20                                  25                                  30

Glu Val Ala Gln Ala Leu Glu Leu Asn Pro Asp His Leu Arg Lys Trp  
                   35                                  40                                  45

Ile Arg Leu Tyr Lys Gln Glu Leu Gln Gly Ile Glu Pro Ala Gly Asn  
                   50                                  55                                  60

Ala Ile Thr Pro Glu Gln Arg Glu Ile Gln Gln Leu Lys Ala Gln Ile  
   65                                  70                                  75                                  80

Lys Arg Val Glu Met Glu Lys Glu Ile Leu Lys Gln Ala Ala Val Leu  
                   85                                  90                                  95

Met Ser Glu Ile Pro Gly Lys Leu Ser Arg  
                   100                                  105

&lt;210&gt; 49

&lt;211&gt; 27

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Oligonucleotide

&lt;400&gt; 49

tgctctagag ccattactca gaatggg

27

<210> 50  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 50  
cgcgagctcg acgactgaat gatccc 26

<210> 51  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 51  
tcccccggtt actgcagcac toaacc 26

<210> 52  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 52  
gatccccgga ccaactgaaat gcgtgc 26

<210> 53  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 53  
tcgtctagag atgatggtga tggagcg 27

<210> 54  
<211> 28  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 54  
gaactgcagc caaatactga taccaccc 28

<210> 55  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 55  
gaactgcagg ctaaaacaga agacgcg 27

<210> 56  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 56  
catgcatgca ctccatatga caaccgc 27

<210> 57  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 57  
tcgtctagaa tgaagctgcg catgagg 27

<210> 58  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 58  
caactgcagt cgcaaatgc gaactgg 27

<210> 59  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 59  
caactgcaga cgcgaacttt togacgc 27

<210> 60  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 60  
catgcatgcc agtgagccat tgttccc 27

<210> 61  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 61  
tgctctagat acgactctga caggagg 27



<210> 62  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 62  
tcagatatca actaccagca gtttgg 26

<210> 63  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 63  
tcagatatcc ataaagagtg acgtggc 27

<210> 64  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 64  
tgctctagaa aacgtggcaa cagagcg 27

<210> 65  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 65  
tgctctagaa ggcgttgctg atcctg 26

<210> 66  
<211> 28  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 66  
gaactgcagg aaaaggccga gcagactg 28

<210> 67  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 67  
gaactgcagt acagccatgt ttacggt 27

<210> 68  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 68  
catgcatgcg gtgtacgaca gtttgcg 27

<210> 69  
<211> 26  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 69  
tgctctagac acatcatggg cacacc 26

<210> 70  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 70  
gaactgcaga accgtccaca tcaggcg 27

<210> 71  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 71  
gaactgcaga ccttgcttgc cattccg 27

<210> 72  
<211> 27  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence:Oligonucleotide

<400> 72  
catgcatgca taagcgtcga acaggcg 27